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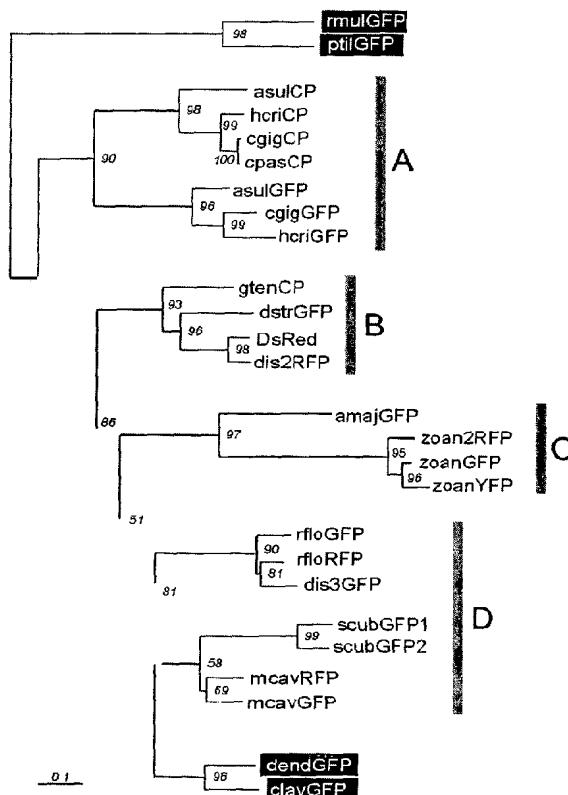
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[Continued on next page]

(54) Title: NOVEL CHROMOPHORES/FLUOROPHORES AND METHODS FOR USING THE SAME



(57) Abstract: Nucleic acid compositions encoding novel chromo/fluoroproteins and mutants thereof, as well as the proteins encoded the same, are provided. The proteins of interest are proteins that are colored and/or fluorescent, where this feature arises from the interaction of two or more residues of the protein. The subject proteins are further characterized in that they are either obtained from non-bioluminescent Cnidarian, e.g., Anthozoan, species or are obtained from Anthozoan non-Pennatulacean (sea pen) species. Specific proteins of interest include the following specific proteins: heriGFP; dendGFP; zoanRFP; scubGFP1; scubGFP2; rfloRFP; rfloGFP; mcavRFP; mcavGFP; cgigGFP; afraGFP; rfloGFP2; mcavGFP2; and mannFP. Also of interest are proteins that are substantially similar to, or mutants of, the above specific proteins. Also provided are fragments of the nucleic acids and the peptides encoded thereby, as well as antibodies to the subject proteins and transgenic cells and organisms. The subject protein and nucleic acid compositions find use in a variety of different applications. Finally, kits for use in such applications, e.g., that include the subject nucleic acid compositions, are provided.

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**NOVEL CHROMOPHORES/FLUOROPHORES AND
METHODS FOR USING THE SAME
CROSS-REFERENCE TO RELATED APPLICATIONS**

Pursuant to 35 U.S.C. § 119 (e), this application claims priority to the filing
5 date of United States Provisional Patent Application Serial No. 60/332,980 filed
November 13, 2001; the disclosure of which is herein incorporated by reference.

INTRODUCTION

Field of the Invention

10 The field of this invention is chromoproteins and fluorescent proteins.

Background of the Invention

Labeling is a tool for marking a protein, cell, or organism of interest and
plays a prominent role in many biochemistry, molecular biology and medical
diagnostic applications. A variety of different labels have been developed,
15 including radiolabels, chromolabels, fluorescent labels, chemiluminescent labels,
etc. However, there is continued interest in the development of new labels. Of
particular interest is the development of new protein labels, including chromo-
and/or fluorescent protein labels.

Relevant Literature

20 U.S. Patents of interest include: 6,066,476; 6,020,192; 5,985,577;
5,976,796; 5,968,750; 5,968,738; 5,958,713; 5,919,445; 5,874,304; and
5,491,084. International Patent Publications of interest include: WO 00/46233;
WO 99/49019; and DE 197 18 640 A. Also of interest are: Anderluh et al.,
Biochemical and Biophysical Research Communications (1996) 220:437-442;
25 Dove et al., Biological Bulletin (1995) 189:288-297; Fradkov et al., FEBS Lett.
(2000) 479(3):127-30; Gurskaya et al., FEBS Lett., (2001) 507(1):16-20; Gurskaya
et al., BMC Biochem. (2001) 2:6; Lukyanov, K., et al (2000) J Biol Chemistry
275(34):25879-25882; Macek et al., Eur. J. Biochem. (1995) 234:329-335;
Martynov et al., J Biol Chem. (2001) 276:21012-6; Matz, M.V., et al. (1999) Nature
30 Biotechnol., 17:969-973; Terskikh et al., Science (2000) 290:1585-8; Tsien, Annual
Rev. of Biochemistry (1998) 67:509-544; Tsien, Nat. Biotech. (1999) 17:956-957;
Ward et al., J. Biol. Chem. (1979) 254:781-788; Wiedermann et al., Jarhrestagung

der Deutschen Gesellschaft für Tropenökologie-gto. Ulm. 17-19.02.1999. Poster P-4.20; Yarbrough et al., Proc Natl Acad Sci U S A (2001) 98:462-7.

SUMMARY OF THE INVENTION

5 Nucleic acid compositions encoding novel chromo/fluoroproteins and mutants thereof, as well as the proteins encoded the same, are provided. The proteins of interest are proteins that are colored and/or fluorescent, where this feature arises from the interaction of two or more residues of the protein. The subject proteins are further characterized in that they are either obtained from non-
10 bioluminescent Cnidarian, e.g., Anthozoan, species or are obtained from Anthozoan non-Pennatulacean (sea pen) species. Specific proteins of interest include the following specific proteins: (1) Green fluorescent protein from *Heteractis crispa* (hcriGFP); (2) Green fluorescent protein from *Dendronephthya* sp. (dendGFP); (3) Red fluorescent protein from *Zoanthus* sp. (zoanRFP); (4)
15 Green fluorescent protein from *Scolymia cubensis* (scubGFP1); (5) Green fluorescent protein from *Scolymia cubensis* (scubGFP2); (6) Red fluorescent protein from *Ricordea florida* (rfloRFP); (7) Green fluorescent protein from *Ricordea florida* (rfloGFP); (8) Red fluorescent protein from *Montastraea cavernosa* (mcavRFP); (9) Green fluorescent protein from *Montastraea cavernosa* (mcavGFP); (10) Green fluorescent protein from *Condylactis gigantea* (cgigGFP);
20 (11) Green fluorescent protein from *Agaricia fragilis* (afraGFP); (12) Green fluorescent protein from *Ricordea florida* (rfloGFP2); (13) Green fluorescent protein from *Montastraea cavernosa* (mcavGFP2); and (14) Green fluorescent protein homolog from *Montastraea annularis* (mannFP). Also of interest are
25 proteins that are substantially similar to, or mutants of, the above specific proteins. Also provided are fragments of the nucleic acids and the peptides encoded thereby, as well as antibodies to the subject proteins and transgenic cells and organisms. The subject protein and nucleic acid compositions find use in a variety of different applications. Finally, kits for use in such applications, e.g., that include
30 the subject nucleic acid compositions, are provided.

BRIEF DESCRIPTION OF THE FIGURES

Figure 1. Changes of emission spectra during maturation of the new red-emitters: zoan2RFP (A, B), mcavRFP (C, D) and rfloRFP (E, F). The excitation

wavelength is given within each graph. Horizontal axis is wavelength in nanometers, vertical axis is fluorescence intensity. Maturation stages: A, C, E – early; B, D, F – late (see Methods for details). All the three proteins exhibit “timer” phenotype (green emission at first and red emission arising later). Note that 5 zoan2RFP matures significantly faster than mcavRFP and rfloRFP: even at the “early” stage the red emission peak is very pronounced, and by the “late” stage the protein converts into red completely. In contrast, mcavRFP and rfloRFP fail to undergo such a complete maturation.

Figure 2. Details on excitation spectra of mcavRFP (A, B) and rfloRFP (C, 10 D). Wavelengths at which the emission was monitored are given within the graphs. A, C: excitation spectra of the green emission band in the immature protein, lacking the red emission; B, D: excitation spectra of the red emission band in more mature form. Horizontal axis is wavelength in nanometers, vertical axis is fluorescence intensity. Note that in both proteins, the major excitation peaks for 15 immature green and mature red forms are virtually identical to each other.

Figure 3. The maximum-likelihood phylogenetic tree for the current dataset 20 of anthozoan GFP-like proteins. Numbers at nodes denote the quartet-puzzling support values (1000 puzzling attempts). Proteins from Alcyonaria sub-class, which were considered outgroups, are labeled in white on black. The “stem” of the tree (thick gray line), joining two rooting groups, putatively reflects the diversity of GFP-like proteins before the separation of Alcyonaria and Zoantharia sub-classes. Gray bars marked A, B, C and D denote four distinct clades of GFP-like proteins found in Zoantharia. Scale bar: 0.1 replacements/site.

Figure 4. Summary of spectral features and chromophore structures in the 25 family of GFP-like proteins. Note that this paper uses different names for GFP-like proteins than proposed in original publications (the original names, where available, are given in brackets in the first column; see text for nomenclature details).

Figure 5. Summary of the major clades of GFP-like proteins from sub-class 30 Zoantharia.

Figure 6. Excitation (solid lines) and emission (dotted lines) spectra for the GFP-like proteins reported in this paper. The wavelengths at which the excitation or emission curves were taken are given in the legend to each graph. Horizontal

axis is wavelength in nanometers, vertical axis is fluorescence intensity. The graphs for the two new orange-red proteins are boxed.

Figure 7. Alignment of the currently cloned and spectroscopically characterized GFP-like proteins. Numeration above the alignment is according to 5 GFP from *Aequorea victoria*.

Figure 8 provides the nucleotide and amino acid sequence of wild type *Heteractis crispa* hcriGFP. (SEQ ID NO:01 & 02)

Figure 9 provides the nucleotide and amino acid sequence of wild type *Dendronephthya* sp. dendGFP. (SEQ ID NO:03 & 04)

10 Figure 10 provides the nucleotide and amino acid sequence of wild type *Zoanthus* sp. zoanRFP. (SEQ ID NO:05 & 06)

Figure 11 provides the nucleotide and amino acid sequence of wild type *Scolymia cubensis* scubGFP1. (SEQ ID NO:07 & 08)

15 Figure 12 provides the nucleotide and amino acid sequence of wild type *Scolymia cubensis* scubGFP2. (SEQ ID NO:09 & 10)

Figure 13 provides the nucleotide and amino acid sequence of wild type *Ricordea florida* rfloRFP. (SEQ ID NO:11 & 12)

Figure 14 provides the nucleotide and amino acid sequence of wild type *Ricordea florida* rfloGFP. (SEQ ID NO:13 & 14)

20 Figure 15 provides the nucleotide and amino acid sequence of wild type *Montastraea cavernosa* mcavRFP. (SEQ ID NO:15 & 16)

Figure 16 provides the nucleotide and amino acid sequence of wild type *Montastraea cavernosa* mcavGFP. (SEQ ID NO:17 & 18)

25 Figure 17 provides the nucleotide and amino acid sequence of wild type *Condylactis gigantea* cgigGFP. (SEQ ID NO: 19 & 20).

Figure 18 provides the nucleotide and amino acid sequence of wild type *Agaricia fragilis* afraGFP. (SEQ ID NO: 21& 22).

Figure 19 provides the nucleotide and amino acid sequence of wild type *Ricordea florida* rfloGFP2. (SEQ ID NO: 23& 24).

30 Figure 20 provides the nucleotide and amino acid sequence of wild type *Montastraea cavernosa* mcavGFP2. (SEQ ID NO: 25& 26).

Figure 21 provides the nucleotide and amino acid sequence of wild type *Montastraea annularis* mannFP. (SEQ ID NO: 27& 28).

FEATURES OF THE INVENTION

The subject invention provides a nucleic acid present in other than its natural environment, wherein the nucleic acid encodes a chromo- or fluorescent protein and is from a non-bioluminescent Cnidarian species. In certain 5 embodiments, the non-bioluminescent Cnidarian species is an Anthozoan species. In certain embodiments, the nucleic acid is isolated. In certain embodiments, the nucleic acid is present in other than its natural environment, where the nucleic acid encodes an Anthozoan chromo- or fluorescent protein and is from a non- 10 Pennatulacean Anthozoan species. In certain embodiments, the nucleic acid has a sequence of residues that is substantially the same as or identical to a nucleotide sequence of at least 10 residues in length of SEQ ID NOS:01, 03, 05, 07, 09, 11, 13, 15, 17; 19; 21; 23; 25; and 27. In certain embodiments, the nucleic acid has a sequence similarity of at least about 60% with a sequence of at least 10 residues 15 in length selected from the group of sequences consisting of SEQ ID NOS:01, 03, 05, 07, 09, 11, 13, 15, 17; 19; 21; 23; 25; and 27. In certain embodiments, the nucleic acid encodes a chromo and/or fluorescent protein that is either: (a) from a 20 non-bioluminescent Cnidarian species; or (b) from a non- Pennatulacean Anthozoan species. In certain embodiments, the nucleic acid encodes a protein that has an amino acid sequence selected from the group consisting of: SEQ ID NOS: 02; 04; 06; 08; 10; 12; 14; 16; 18; 20; 22; 24; 26; and 28. In certain 25 embodiments, the nucleic acid encodes a mutant protein of a chromo and/or fluorescent protein that is either: (a) from a non-bioluminescent Cnidarian species; or (b) from a non- Pennatulacean Anthozoan species; where in certain embodiments the mutant protein comprises at least one point mutation as compared to its wild type protein; and in other embodiments the mutant protein comprises at least one deletion mutation as compared to its wild type protein.

Also provided are fragments of the provided nucleic acids. Also provided are isolated nucleic acids or mimetics thereof that hybridize under stringent 30 conditions to the provided nucleic acids. Also provided are constructs comprising a vector and a nucleic acid of the present invention. Also provided are expression cassettes that include: (a) a transcriptional initiation region functional in an expression host; (b) a nucleic acid of the present invention; and (c) a transcriptional termination region functional in said expression host. Also provided

are cells, or the progeny thereof, comprising an expression cassette of the present invention as part of an extrachromosomal element or integrated into the genome of a host cell as a result of introduction of said expression cassette into said host cell.

5 Also provided are methods of producing a chromo and/or fluorescent protein that include: growing a cell of the present invention, whereby said protein is expressed; and isolating said protein substantially free of other proteins.

Also provided are proteins or fragments thereof encoded by a nucleic acid of the present invention.

10 Also provided are antibodies binding specifically to a protein of the present invention.

Also provided are transgenic cells or the progeny thereof that include a transgene selected that includes a nucleic acid of the present invention.

15 Also provided are transgenic organisms that include a transgene that includes a nucleic acid of the present invention.

Also provided are applications that employ a chromo- or fluorescent protein of the present invention.

Also provided are applications that employ a nucleic acid encoding a chromo- or fluorescent protein of the present invention.

20 Also provided are kits that include a nucleic acid according the subject invention and instructions for using said nucleic acid.

DEFINITIONS

In accordance with the present invention there may be employed conventional molecular biology, microbiology, and recombinant DNA techniques within the skill of the art. Such techniques are explained fully in the literature. See, e.g., Maniatis, Fritsch & Sambrook, "Molecular Cloning: A Laboratory Manual" (1982); "DNA Cloning: A Practical Approach," Volumes I and II (D.N. Glover ed. 1985); "Oligonucleotide Synthesis" (M.J. Gait ed. 1984); "Nucleic Acid Hybridization" (B.D. Hames & S.J. Higgins eds. (1985)); "Transcription and Translation" (B.D. Hames & S.J. Higgins eds. (1984)); "Animal Cell Culture" (R.I. Freshney, ed. (1986)); "Immobilized Cells and Enzymes" (IRL Press, (1986)); B. Perbal, "A Practical Guide To Molecular Cloning" (1984).

A "vector" is a replicon, such as plasmid, phage or cosmid, to which another DNA segment may be attached so as to bring about the replication of the attached segment.

A "DNA molecule" refers to the polymeric form of deoxyribonucleotides (adenine, guanine, thymine, or cytosine) in either single stranded form or a double-stranded helix. This term refers only to the primary and secondary structure of the molecule, and does not limit it to any particular tertiary forms. Thus, this term includes double-stranded DNA found, *inter alia*, in linear DNA molecules (e.g., restriction fragments), viruses, plasmids, and chromosomes.

A DNA "coding sequence" is a DNA sequence which is transcribed and translated into a polypeptide *in vivo* when placed under the control of appropriate regulatory sequences. The boundaries of the coding sequence are determined by a start codon at the 5' (amino) terminus and a translation stop codon at the 3' (carboxyl) terminus. A coding sequence can include, but is not limited to, prokaryotic sequences, cDNA from eukaryotic mRNA, genomic DNA sequences from eukaryotic (e.g., mammalian) DNA, and synthetic DNA sequences. A polyadenylation signal and transcription termination sequence may be located 3' to the coding sequence.

As used herein, the term "hybridization" refers to the process of association of two nucleic acid strands to form an antiparallel duplex stabilized by means of hydrogen bonding between residues of the opposite nucleic acid strands.

The term "oligonucleotide" refers to a short (under 100 bases in length) nucleic acid molecule.

"DNA regulatory sequences", as used herein, are transcriptional and translational control sequences, such as promoters, enhancers, polyadenylation signals, terminators, and the like, that provide for and/or regulate expression of a coding sequence in a host cell.

A "promoter sequence" is a DNA regulatory region capable of binding RNA polymerase in a cell and initiating transcription of a downstream (3' direction) coding sequence. For purposes of defining the present invention, the promoter sequence is bounded at its 3' terminus by the transcription initiation site and extends upstream (5' direction) to include the minimum number of bases or elements necessary to initiate transcription at levels detectable above background. Within the promoter sequence will be found a transcription initiation site, as well as

protein binding domains responsible for the binding of RNA polymerase. Eukaryotic promoters will often, but not always, contain "TATA" boxes and "CAT" boxes. Various promoters, including inducible promoters, may be used to drive the various vectors of the present invention.

5 As used herein, the terms "restriction endonucleases" and "restriction enzymes" refer to bacterial enzymes, each of which cut double-stranded DNA at or near a specific nucleotide sequence.

10 A cell has been "transformed" or "transfected" by exogenous or heterologous DNA when such DNA has been introduced inside the cell. The transforming DNA may or may not be integrated (covalently linked) into the genome of the cell. In prokaryotes, yeast, and mammalian cells for example, the transforming DNA may be maintained on an episomal element such as a plasmid. With respect to eukaryotic cells, a stably transformed cell is one in which the transforming DNA has become integrated into a chromosome so that it is inherited 15 by daughter cells through chromosome replication. This stability is demonstrated by the ability of the eukaryotic cell to establish cell lines or clones comprised of a population of daughter cells containing the transforming DNA. A "clone" is a population of cells derived from a single cell or common ancestor by mitosis. A "cell line" is a clone of a primary cell that is capable of stable growth *in vitro* for 20 many generations.

25 A "heterologous" region of the DNA construct is an identifiable segment of DNA within a larger DNA molecule that is not found in association with the larger molecule in nature. Thus, when the heterologous region encodes a mammalian genomic DNA in the genome of the source organism. In another example, heterologous DNA includes coding sequence in a construct where portions of genes from two different sources have been brought together so as to produce a fusion protein product. Allelic variations or naturally-occurring mutational events do not give rise to a heterologous region of DNA as defined herein.

30 As used herein, the term "reporter gene" refers to a coding sequence attached to heterologous promoter or enhancer elements and whose product may be assayed easily and quantifiably when the construct is introduced into tissues or cells.

The amino acids described herein are preferred to be in the "L" isomeric form. The amino acid sequences are given in one-letter code (A: alanine; C: cysteine; D: aspartic acid; E: glutamic acid; F: phenylalanine; G: glycine; H: histidine; I: isoleucine; K: lysine; L: leucine; M: methionine; N: asparagine; P: 5 proline; Q: glutamine; R: arginine; S: serine; T: threonine; V: valine; W: tryptophan; Y: tyrosine; X: any residue). NH₂ refers to the free amino group present at the amino terminus of a polypeptide. COOH refers to the free carboxy group present at the carboxy terminus of a polypeptide. In keeping with standard polypeptide nomenclature, *J Biol. Chem.*, 243 (1969), 3552-59 is used.

10 The term "immunologically active" defines the capability of the natural, recombinant or synthetic chromo/fluorescent protein, or any oligopeptide thereof, to induce a specific immune response in appropriate animals or cells and to bind with specific antibodies. As used herein, "antigenic amino acid sequence" means an amino acid sequence that, either alone or in association with a carrier 15 molecule, can elicit an antibody response in a mammal. The term "specific binding," in the context of antibody binding to an antigen, is a term well understood in the art and refers to binding of an antibody to the antigen to which the antibody was raised, but not other, unrelated antigens.

20 As used herein the term "isolated" is meant to describe a polynucleotide, a polypeptide, an antibody, or a host cell that is in an environment different from that in which the polynucleotide, the polypeptide, the antibody, or the host cell naturally occurs.

25 Bioluminescence (BL) is defined as emission of light by living organisms that is well visible in the dark and affects visual behavior of animals (See e.g., Harvey, E. N. (1952). *Bioluminescence*. New York: Academic Press; Hastings, J. W. (1995). Bioluminescence. In: *Cell Physiology* (ed. by N. Speralakis). pp. 651-681. New York: Academic Press.; Wilson, T. and Hastings, J. W. (1998). Bioluminescence. *Annu Rev Cell Dev Biol* **14**, 197-230.). Bioluminescence does not include so-called ultra-weak light emission, which can be detected in virtually 30 all living structures using sensitive luminometric equipment (Murphy, M. E. and Sies, H. (1990). Visible-range low-level chemiluminescence in biological systems. *Meth. Enzymol.* **186**, 595-610; Radotic, K, Radenovic, C, Jeremic, M. (1998.) Spontaneous ultra-weak bioluminescence in plants: origin, mechanisms and

properties. *Gen Physiol Biophys* **17**, 289-308), and from weak light emission which most probably does not play any ecological role, such as the glowing of bamboo growth cone (Totsune, H., Nakano, M., Inaba, H. (1993). Chemiluminescence from bamboo shoot cut. *Biochem. Biophys. Res Comm.* **194**, 1025-1029) or emission of light during fertilization of animal eggs (Klebanoff, S. J., Froeder, C. A., Eddy, E. M., Shapiro, B. M. (1979). Metabolic similarities between fertilization and phagocytosis. Conservation of peroxidatic mechanism. *J. Exp. Med.* **149**, 938-953; Schomer, B. and Epel, D. (1998). Redox changes during fertilization and maturation of marine invertebrate eggs. *Dev Biol* **203**, 1-11).

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DESCRIPTION OF THE SPECIFIC EMBODIMENTS

Nucleic acid compositions encoding novel chromo/fluoroproteins and mutants thereof, as well as the proteins encoded the same, are provided. The proteins of interest are proteins that are colored and/or fluorescent, where this feature arises from the interaction of two or more residues of the protein. The subject proteins are further characterized in that they are either obtained from non-bioluminescent Cnidarian, e.g., Anthozoan, species or are obtained from non-Pennatulacean (sea pen) Anthozoan species. Specific proteins of interest include the following specific proteins: (1) Green fluorescent protein from *Heteractis crispa* (hcriGFP); (2) Green fluorescent protein from *Dendronephthya* sp. (dendGFP); (3) Red fluorescent protein from *Zoanthus* sp. (zoanRFP); (4) Green fluorescent protein from *Scolymia cubensis* (scubGFP1); (5) Green fluorescent protein from *Scolymia cubensis* (scubGFP2); (6) Red fluorescent protein from *Ricordea florida* (rfloRFP); (7) Green fluorescent protein from *Ricordea florida* (rfloGFP); (8) Red fluorescent protein from *Montastraea cavernosa* (mcavRFP); (9) Green fluorescent protein from *Montastraea cavernosa* (mcavGFP); (10) Green fluorescent protein from *Condylactis gigantea* (cgigGFP); (11) Green fluorescent protein from *Agaricia fragilis* (afraGFP); (12) Green fluorescent protein from *Ricordea florida* (rfloGFP2); (13) Green fluorescent protein from *Montastraea cavernosa* (mcavGFP2); and (14) Green fluorescent protein homolog from *Montastraea annularis* (mannFP). Also of interest are proteins that are substantially similar to, or mutants of, the above specific proteins. Also provided are fragments of the nucleic acids and the peptides encoded thereby, as well as antibodies to the subject proteins, and

transgenic cells and organisms that include the subject nucleic acid/protein compositions. The subject protein and nucleic acid compositions find use in a variety of different applications. Finally, kits for use in such applications, e.g., that include the subject nucleic acid compositions, are provided.

5

Before the subject invention is described further, it is to be understood that the invention is not limited to the particular embodiments of the invention described below, as variations of the particular embodiments may be made and still fall within the scope of the appended claims. It is also to be understood that 10 the terminology employed is for the purpose of describing particular embodiments, and is not intended to be limiting. Instead, the scope of the present invention will be established by the appended claims.

In this specification and the appended claims, the singular forms "a," "an" 15 and "the" include plural reference unless the context clearly dictates otherwise. Unless defined otherwise, all technical and scientific terms used herein have the same meaning as commonly understood to one of ordinary skill in the art to which this invention belongs.

20 Where a range of values is provided, it is understood that each intervening value, to the tenth of the unit of the lower limit unless the context clearly dictates otherwise, between the upper and lower limit of that range, and any other stated or intervening value in that stated range, is encompassed within the invention. The upper and lower limits of these smaller ranges may independently be included in 25 the smaller ranges, and are also encompassed within the invention, subject to any specifically excluded limit in the stated range. Where the stated range includes one or both of the limits, ranges excluding either or both of those included limits are also included in the invention.

30 Unless defined otherwise, all technical and scientific terms used herein have the same meaning as commonly understood to one of ordinary skill in the art to which this invention belongs. Although any methods, devices and materials similar or equivalent to those described herein can be used in the practice or

testing of the invention, the preferred methods, devices and materials are now described.

5 All publications mentioned herein are incorporated herein by reference for the purpose of describing and disclosing the cell lines, vectors, methodologies and other invention components that are described in the publications which might be used in connection with the presently described invention.

10 In further describing the subject invention, the subject nucleic acid compositions will be described first, followed by a discussion of the subject protein compositions, antibody compositions and transgenic cells/organisms. Next a review of representative methods in which the subject proteins find use is provided.

15 **NUCLEIC ACID COMPOSITIONS**

As summarized above, the subject invention provides nucleic acid compositions encoding chromo- and fluoroproteins and mutants thereof, as well as fragments and homologues of these proteins. By chromo and/or fluorescent protein is meant a protein that is colored, i.e., is pigmented, where the protein may or may not be fluorescent, e.g., it may exhibit low, medium or high fluorescence upon irradiation with light of an excitation wavelength. In any event, the subject proteins of interest are those in which the colored characteristic, i.e., the chromo and/or fluorescent characteristic, is one that arises from the interaction of two or 25 more residues of the protein, and not from a single residue, more specifically a single side chain of a single residue, of the protein. As such, fluorescent proteins of the subject invention do not include proteins that exhibit fluorescence only from residues that act by themselves as intrinsic fluors, i.e., tryptophan, tyrosine and phenylalanine. As such, the fluorescent proteins of the subject invention are 30 fluorescent proteins whose fluorescence arises from some structure in the protein that is other than the above specified single residues, e.g., it arises from an interaction of two or more residues.

By nucleic acid composition is meant a composition comprising a sequence of DNA having an open reading frame that encodes a chromo/fluoro polypeptide of

the subject invention, i.e., a chromo/fluoroprotein gene, and is capable, under appropriate conditions, of being expressed as a chromo/fluoro protein according to the subject invention. Also encompassed in this term are nucleic acids that are homologous, substantially similar or identical to the nucleic acids of the present invention. Thus, the subject invention provides genes and coding sequences thereof encoding the proteins of the subject invention, as well as homologs thereof. The subject nucleic acids are present in other than their natural environment, e.g., they are isolated, present in enriched amounts, etc., from their naturally occurring environment, e.g., the organism from which they are obtained.

The nucleic acids are further characterized in that they encode proteins that are either from: (1) non-bioluminescent species, often non-bioluminescent Cnidarian species, e.g., non-bioluminescent Anthozoan species; or (2) from Anthozoan species that are not Pennatulacean species, i.e., that are not sea pens. As such, the nucleic acids may encode proteins from bioluminescent Anthozoan species, so long as these species are not Pennatulacean species, e.g., that are not Renillan or Ptilosarcan species. Specific nucleic acids of interest are those that encode the following specific proteins: (1) Green fluorescent protein from *Heteractis crispa* (hcriGFP) (Genbank Accession No. AF420592); (2) Green fluorescent protein from *Dendronephthya* sp. (dendGFP) (Genbank Accession No. AF420591); (3) Red fluorescent protein from *Zoanthus* sp. (zoanRFP) (Genbank Accession No. AY059642); (4) Green fluorescent protein from *Scolymia cubensis* (scubGFP1) (Genbank Accession No. AY037767); (5) Green fluorescent protein from *Scolymia cubensis* (scubGFP2) (Genbank Accession No. AY037771); (6) Red fluorescent protein from *Ricordea florida* (rfloRFP) (Genbank Accession No. AY037773); (7) Green fluorescent protein from *Ricordea florida* (rfloGFP) (Genbank Accession No. AY037772); (8) Red fluorescent protein from *Montastraea cavernosa* (mcavRFP) (Genbank Accession No. AY037770); (9) Green fluorescent protein from *Montastraea cavernosa* (mcavGFP) (Genbank Accession No. AY037769); (10) Green fluorescent protein from *Condylactis gigantea* (cgigGFP) (Genbank Accession No. AY03776); (11) Green fluorescent protein from *Agaricia fragilis* (afraGFP); (12) Green fluorescent protein from *Ricordea florida* (rfloGFP2); (13) Green fluorescent protein from *Montastraea cavernosa* (mcavGFP2); and (14) Green fluorescent protein homolog from

Montastraea annularis (mannFP). Also of interest are derived from, or are mutants, homologues of, the above specific nucleic acids.

In addition to the above described specific nucleic acid compositions, also of interest are homologues of the above sequences. With respect to homologues 5 of the subject nucleic acids, the source of homologous genes may be any species of plant or animal or the sequence may be wholly or partially synthetic. In certain embodiments, sequence similarity between homologues is at least about 20%, sometimes at least about 25 %, and may be 30 %, 35%, 40%, 50%, 60%, 70% or higher, including 75%, 80%, 85%, 90% and 95% or higher. Sequence similarity is 10 calculated based on a reference sequence, which may be a subset of a larger sequence, such as a conserved motif, coding region, flanking region, etc. A reference sequence will usually be at least about 18 nt long, more usually at least about 30 nt long, and may extend to the complete sequence that is being compared. Algorithms for sequence analysis are known in the art, such as 15 BLAST, described in Altschul *et al.* (1990), *J. Mol. Biol.* 215:403-10 (using default settings, i.e. parameters $w=4$ and $T=17$). The sequences provided herein are essential for recognizing related and homologous nucleic acids in database 20 searches. Of particular interest in certain embodiments are nucleic acids of substantially the same length as the nucleic acid identified as SEQ ID NOS: 01, 03, 05, 07, 09, 11, 13, 15, 17, 19, 21, 23, 25 or 27, where by substantially the same length is meant that any difference in length does not exceed about 20 number %, usually does not exceed about 10 number % and more usually does not exceed about 5 number %; and have sequence identity to any of these 25 sequences of at least about 90%, usually at least about 95% and more usually at least about 99% over the entire length of the nucleic acid. In many embodiments, the nucleic acids have a sequence that is substantially similar (i.e. the same as) or identical to the sequences of SEQ ID NOS: 01, 03, 05, 07, 09, 11, 13, 15, 17, 21, 23, 25, 27. By substantially similar is meant that sequence identity will generally be 30 at least about 60%, usually at least about 75% and often at least about 80, 85, 90, or even 95%.

Also provided are nucleic acids that encode the proteins encoded by the above described nucleic acids, but differ in sequence from the above described nucleic acids due to the degeneracy of the genetic code.

Also provided are nucleic acids that hybridize to the above described nucleic acid under stringent conditions. An example of stringent hybridization conditions is hybridization at 50°C or higher and 0.1×SSC (15 mM sodium chloride/1.5 mM sodium citrate). Another example of stringent hybridization conditions is overnight incubation at 42°C in a solution: 50 % formamide, 5 × SSC (150 mM NaCl, 15 mM trisodium citrate), 50 mM sodium phosphate (pH7.6), 5 × Denhardt's solution, 10% dextran sulfate, and 20 µg/ml denatured, sheared salmon sperm DNA, followed by washing the filters in 0.1 × SSC at about 65°C. Stringent hybridization conditions are hybridization conditions that are at least as stringent as the above representative conditions, where conditions are considered to be at least as stringent if they are at least about 80% as stringent, typically at least about 90% as stringent as the above specific stringent conditions. Other stringent hybridization conditions are known in the art and may also be employed to identify nucleic acids of this particular embodiment of the invention.

Nucleic acids encoding mutants of the proteins of the invention are also provided. Mutant nucleic acids can be generated by random mutagenesis or targeted mutagenesis, using well-known techniques which are routine in the art. In some embodiments, chromo- or fluorescent proteins encoded by nucleic acids encoding homologues or mutants have the same fluorescent properties as the wild-type fluorescent protein. In other embodiments, homologue or mutant nucleic acids encode chromo- or fluorescent proteins with altered spectral properties, as described in more detail herein.

One category of mutant that is of particular interest is the non-aggregating mutant. In many embodiments, the non-aggregating mutant differs from the wild type sequence by a mutation in the N-terminus that modulates the charges appearing on side groups of the N-terminus residues, e.g., to reverse or neutralize the charge, in a manner sufficient to produce a non-aggregating mutant of the naturally occurring protein or mutant, where a particular protein is considered to be non-aggregating if it is determined be non-aggregating using the assay reported in U.S. Patent Application serial no. 60/270,983, the disclosure of which is herein incorporated by reference.

Another category of mutant of particular interest is the modulated oligomerization mutant. A mutant is considered to be a modulated oligomerization

mutant if its oligomerization properties are different as compared to the wild type protein. For example, if a particular mutant oligomerizes to a greater or lesser extent than the wild type, it is considered to be an oligomerization mutant. Of particular interest are oligomerization mutants that do not oligomerize, i.e., are 5 monomers under physiological (e.g., intracellular) conditions, or oligomerize to a lesser extent than the wild type, e.g., are dimers or trimers under intracellular conditions.

Nucleic acids of the subject invention may be cDNA or genomic DNA or a fragment thereof. In certain embodiments, the nucleic acids of the subject 10 invention include one or more of the open reading frames encoding specific fluorescent proteins and polypeptides, and introns, as well as adjacent 5' and 3' non-coding nucleotide sequences involved in the regulation of expression, up to about 20 kb beyond the coding region, but possibly further in either direction. The subject nucleic acids may be introduced into an appropriate vector for 15 extrachromosomal maintenance or for integration into a host genome, as described in greater detail below.

The term "cDNA" as used herein is intended to include all nucleic acids that share the arrangement of sequence elements found in native mature mRNA species, where sequence elements are exons and 5' and 3' non-coding regions. 20 Normally mRNA species have contiguous exons, with the intervening introns, when present, being removed by nuclear RNA splicing, to create a continuous open reading frame encoding the protein.

A genomic sequence of interest comprises the nucleic acid present 25 between the initiation codon and the stop codon, as defined in the listed sequences, including all of the introns that are normally present in a native chromosome. It may further include 5' and 3' un-translated regions found in the mature mRNA. It may further include specific transcriptional and translational regulatory sequences, such as promoters, enhancers, etc., including about 1 kb, but possibly more, of flanking genomic DNA at either the 5' or 3' end of the 30 transcribed region. The genomic DNA may be isolated as a fragment of 100 kbp or smaller; and substantially free of flanking chromosomal sequence. The genomic DNA flanking the coding region, either 3' or 5', or internal regulatory

sequences as sometimes found in introns, contains sequences required for proper tissue and stage specific expression.

The nucleic acid compositions of the subject invention may encode all or a part of the subject proteins. Double or single stranded fragments may be obtained from the DNA sequence by chemically synthesizing oligonucleotides in accordance with conventional methods, by restriction enzyme digestion, by PCR amplification, etc. For the most part, DNA fragments will be of at least about 15 nt, usually at least about 18 nt or about 25 nt, and may be at least about 50 nt. In some embodiments, the subject nucleic acid molecules may be about 100 nt, about 200 nt, about 300 nt, about 400 nt, about 500 nt, about 600 nt, about 700 nt, or about 720 nt in length. The subject nucleic acids may encode fragments of the subject proteins or the full-length proteins, e.g., the subject nucleic acids may encode polypeptides of about 25 aa, about 50 aa, about 75 aa, about 100 aa, about 125 aa, about 150 aa, about 200 aa, about 210 aa, about 220 aa, about 230 aa, or about 240 aa, up to the entire protein.

The subject nucleic acids are isolated and obtained in substantial purity, generally as other than an intact chromosome. Usually, the DNA will be obtained substantially free of other nucleic acid sequences that do not include a nucleic acid of the subject invention or fragment thereof, generally being at least about 50%, usually at least about 90% pure and are typically "recombinant", i.e. flanked by one or more nucleotides with which it is not normally associated on a naturally occurring chromosome.

The subject polynucleotides (e.g., a polynucleotide having a sequence of SEQ ID NOS: 01, 03, 05, 07, 09, 11, 13, 15, 17, 19, 21, 23, 25, 27 etc.), the corresponding cDNA, the full-length gene and constructs of the subject polynucleotides are provided. These molecules can be generated synthetically by a number of different protocols known to those of skill in the art. Appropriate polynucleotide constructs are purified using standard recombinant DNA techniques as described in, for example, Sambrook *et al.*, *Molecular Cloning: A Laboratory Manual, 2nd Ed.*, (1989) Cold Spring Harbor Press, Cold Spring Harbor, NY, and under current regulations described in United States Dept. of HHS, National Institute of Health (NIH) Guidelines for Recombinant DNA Research.

Also provided are nucleic acids that encode fusion proteins of the subject proteins, or fragments thereof, which are fused to a second protein, e.g., a degradation sequence, a signal peptide, etc. Fusion proteins may comprise a subject polypeptide, or fragment thereof, and a non-Anthozoan polypeptide ("the fusion partner") fused in-frame at the N-terminus and/or C-terminus of the subject polypeptide. Fusion partners include, but are not limited to, polypeptides that can bind antibody specific to the fusion partner (e.g., epitope tags); antibodies or binding fragments thereof; polypeptides that provide a catalytic function or induce a cellular response; ligands or receptors or mimetics thereof; and the like. In such fusion proteins, the fusion partner is generally not naturally associated with the subject Anthozoan portion of the fusion protein, and is typically not an Anthozoan protein or derivative/fragment thereof, i.e., it is not found in Anthozoan species.

Also provided are constructs comprising the subject nucleic acids inserted into a vector, where such constructs may be used for a number of different applications, including propagation, protein production, etc. Viral and non-viral vectors may be prepared and used, including plasmids. The choice of vector will depend on the type of cell in which propagation is desired and the purpose of propagation. Certain vectors are useful for amplifying and making large amounts of the desired DNA sequence. Other vectors are suitable for expression in cells in culture. Still other vectors are suitable for transfer and expression in cells in a whole animal or person. The choice of appropriate vector is well within the skill of the art. Many such vectors are available commercially. To prepare the constructs, the partial or full-length polynucleotide is inserted into a vector typically by means of DNA ligase attachment to a cleaved restriction enzyme site in the vector.

Alternatively, the desired nucleotide sequence can be inserted by homologous recombination *in vivo*. Typically this is accomplished by attaching regions of homology to the vector on the flanks of the desired nucleotide sequence. Regions of homology are added by ligation of oligonucleotides, or by polymerase chain reaction using primers comprising both the region of homology and a portion of the desired nucleotide sequence, for example.

Also provided are expression cassettes or systems that find use in, among other applications, the synthesis of the subject proteins. For expression, the gene product encoded by a polynucleotide of the invention is expressed in any convenient expression system, including, for example, bacterial, yeast, insect,

amphibian and mammalian systems. Suitable vectors and host cells are described in U.S. Patent No. 5,654,173. In the expression vector, a subject polynucleotide, e.g., as set forth in SEQ ID NOS:01; 03; 05; 07; 09; 11; 13; 15; 17; 19; 21; 23; 25 or 27, is linked to a regulatory sequence as appropriate to obtain 5 the desired expression properties. These regulatory sequences can include promoters (attached either at the 5' end of the sense strand or at the 3' end of the antisense strand), enhancers, terminators, operators, repressors, and inducers. The promoters can be regulated or constitutive. In some situations it may be desirable to use conditionally active promoters, such as tissue-specific or 10 developmental stage-specific promoters. These are linked to the desired nucleotide sequence using the techniques described above for linkage to vectors. Any techniques known in the art can be used. In other words, the expression vector will provide a transcriptional and translational initiation region, which may be inducible or constitutive, where the coding region is operably linked under the 15 transcriptional control of the transcriptional initiation region, and a transcriptional and translational termination region. These control regions may be native to the subject species from which the subject nucleic acid is obtained, or may be derived from exogenous sources.

Expression vectors generally have convenient restriction sites located near 20 the promoter sequence to provide for the insertion of nucleic acid sequences encoding heterologous proteins. A selectable marker operative in the expression host may be present. Expression vectors may be used for, among other things, the production of fusion proteins, as described above.

Expression cassettes may be prepared comprising a transcription initiation 25 region, the gene or fragment thereof, and a transcriptional termination region. Of particular interest is the use of sequences that allow for the expression of functional epitopes or domains, usually at least about 8 amino acids in length, more usually at least about 15 amino acids in length, to about 25 amino acids, and up to the complete open reading frame of the gene. After introduction of the DNA, 30 the cells containing the construct may be selected by means of a selectable marker, the cells expanded and then used for expression.

The above described expression systems may be employed with prokaryotes or eukaryotes in accordance with conventional ways, depending upon the purpose for expression. For large scale production of the protein, a unicellular

organism, such as *E. coli*, *B. subtilis*, *S. cerevisiae*, insect cells in combination with baculovirus vectors, or cells of a higher organism such as vertebrates, e.g. COS 7 cells, HEK 293, CHO, Xenopus Oocytes, etc., may be used as the expression host cells. In some situations, it is desirable to express the gene in eukaryotic cells, 5 where the expressed protein will benefit from native folding and post-translational modifications. Small peptides can also be synthesized in the laboratory. Polypeptides that are subsets of the complete protein sequence may be used to identify and investigate parts of the protein important for function.

Specific expression systems of interest include bacterial, yeast, insect cell 10 and mammalian cell derived expression systems. Representative systems from each of these categories is are provided below:

Bacteria. Expression systems in bacteria include those described in Chang et al., *Nature* (1978) 275:615; Goeddel et al., *Nature* (1979) 281:544; Goeddel et al., *Nucleic Acids Res.* (1980) 8:4057; EP 0 036,776; U.S. Patent No. 4,551,433; 15 DeBoer et al., *Proc. Natl. Acad. Sci. (USA)* (1983) 80:21-25; and Siebenlist et al., *Cell* (1980) 20:269.

Yeast. Expression systems in yeast include those described in Hinnen et al., *Proc. Natl. Acad. Sci. (USA)* (1978) 75:1929; Ito et al., *J. Bacteriol.* (1983) 153:163; Kurtz et al., *Mol. Cell. Biol.* (1986) 6:142; Kunze et al., *J. Basic Microbiol.* 20 (1985) 25:141; Gleeson et al., *J. Gen. Microbiol.* (1986) 132:3459; Roggenkamp et al., *Mol. Gen. Genet.* (1986) 202:302; Das et al., *J. Bacteriol.* (1984) 158:1165; De Louvencourt et al., *J. Bacteriol.* (1983) 154:737; Van den Berg et al., *Bio/Technology* (1990) 8:135; Kunze et al., *J. Basic Microbiol.* (1985) 25:141; Cregg et al., *Mol. Cell. Biol.* (1985) 5:3376; U.S. Patent Nos. 4,837,148 and 25 4,929,555; Beach and Nurse, *Nature* (1981) 300:706; Davidow et al., *Curr. Genet.* (1985) 10:380; Gaillardin et al., *Curr. Genet.* (1985) 10:49; Ballance et al., *Biochem. Biophys. Res. Commun.* (1983) 112:284-289; Tilburn et al., *Gene* (1983) 26:205-221; Yelton et al., *Proc. Natl. Acad. Sci. (USA)* (1984) 81:1470-1474; Kelly and Hynes, *EMBO J.* (1985) 4:475479; EP 0 244,234; and WO 91/00357.

30 Insect Cells. Expression of heterologous genes in insects is accomplished as described in U.S. Patent No. 4,745,051; Friesen et al., "The Regulation of Baculovirus Gene Expression", in: *The Molecular Biology Of Baculoviruses* (1986) (W. Doerfler, ed.); EP 0 127,839; EP 0 155,476; and Vlak et al., *J. Gen. Virol.*

(1988) 69:765-776; Miller *et al.*, *Ann. Rev. Microbiol.* (1988) 42:177; Carbonell *et al.*, *Gene* (1988) 73:409; Maeda *et al.*, *Nature* (1985) 315:592-594; Lebacq-Verheyden *et al.*, *Mol. Cell. Biol.* (1988) 8:3129; Smith *et al.*, *Proc. Natl. Acad. Sci. (USA)* (1985) 82:8844; Miyajima *et al.*, *Gene* (1987) 58:273; and Martin 5 *et al.*, *DNA* (1988) 7:99. Numerous baculoviral strains and variants and corresponding permissive insect host cells from hosts are described in Luckow *et al.*, *Bio/Technology* (1988) 6:47-55, Miller *et al.*, *Generic Engineering* (1986) 8:277-279, and Maeda *et al.*, *Nature* (1985) 315:592-594.

10 Mammalian Cells. Mammalian expression is accomplished as described in Dijkema *et al.*, *EMBO J.* (1985) 4:761, Gorman *et al.*, *Proc. Natl. Acad. Sci. (USA)* (1982) 79:6777, Boshart *et al.*, *Cell* (1985) 41:521 and U.S. Patent No. 4,399,216. Other features of mammalian expression are facilitated as described in Ham and Wallace, *Meth. Enz.* (1979) 58:44, Barnes and Sato, *Anal. Biochem.* (1980) 102:255, U.S. Patent Nos. 4,767,704, 4,657,866, 4,927,762, 4,560,655, WO 15 90/103430, WO 87/00195, and U.S. RE 30,985.

20 When any of the above host cells, or other appropriate host cells or organisms, are used to replicate and/or express the polynucleotides or nucleic acids of the invention, the resulting replicated nucleic acid, RNA, expressed protein or polypeptide, is within the scope of the invention as a product of the host cell or organism. The product is recovered by any appropriate means known in the art.

25 Once the gene corresponding to a selected polynucleotide is identified, its expression can be regulated in the cell to which the gene is native. For example, an endogenous gene of a cell can be regulated by an exogenous regulatory sequence inserted into the genome of the cell at location sufficient to at least enhance expression of the gene in the cell. The regulatory sequence may be designed to integrate into the genome via homologous recombination, as disclosed in U.S. Patent Nos. 5,641,670 and 5,733,761, the disclosures of which are herein incorporated by reference, or may be designed to integrate into the 30 genome via non-homologous recombination, as described in WO 99/15650, the disclosure of which is herein incorporated by reference. As such, also encompassed in the subject invention is the production of the subject proteins without manipulation of the encoding nucleic acid itself, but instead through

integration of a regulatory sequence into the genome of cell that already includes a gene encoding the desired protein, as described in the above incorporated patent documents.

Also provided are homologs of the subject nucleic acids. Homologs are identified by any of a number of methods. A fragment of the provided cDNA may be used as a hybridization probe against a cDNA library from the target organism of interest, where low stringency conditions are used. The probe may be a large fragment, or one or more short degenerate primers. Nucleic acids having sequence similarity are detected by hybridization under low stringency conditions, for example, at 50°C and 6×SSC (0.9 M sodium chloride/0.09 M sodium citrate) and remain bound when subjected to washing at 55°C in 1×SSC (0.15 M sodium chloride/.015 M sodium citrate). Sequence identity may be determined by hybridization under stringent conditions, for example, at 50°C or higher and 0.1×SSC (15 mM sodium chloride/1.5 mM sodium citrate). Nucleic acids having a region of substantial identity to the provided sequences, e.g. allelic variants, genetically altered versions of the gene, etc., bind to the provided sequences under stringent hybridization conditions. By using probes, particularly labeled probes of DNA sequences, one can isolate homologous or related genes.

Also of interest are promoter elements of the subject genomic sequences, where the sequence of the 5' flanking region may be utilized for promoter elements, including enhancer binding sites, e.g., that provide for regulation of expression in cells/tissues where the subject proteins gene are expressed.

Also provided are small DNA fragments of the subject nucleic acids, which fragments are useful as primers for PCR, hybridization screening probes, etc. Larger DNA fragments, *i.e.*, greater than 100 nt are useful for production of the encoded polypeptide, as described in the previous section. For use in geometric amplification reactions, such as geometric PCR, a pair of primers will be used. The exact composition of the primer sequences is not critical to the invention, but for most applications the primers will hybridize to the subject sequence under stringent conditions, as known in the art. It is preferable to choose a pair of primers that will generate an amplification product of at least about 50 nt, preferably at least about 100 nt. Algorithms for the selection of primer sequences are generally known, and are available in commercial software packages.

Amplification primers hybridize to complementary strands of DNA, and will prime towards each other.

The DNA may also be used to identify expression of the gene in a biological specimen. The manner in which one probes cells for the presence of particular 5 nucleotide sequences, as genomic DNA or RNA, is well established in the literature. Briefly, DNA or mRNA is isolated from a cell sample. The mRNA may be amplified by RT-PCR, using reverse transcriptase to form a complementary DNA strand, followed by polymerase chain reaction amplification using primers specific for the subject DNA sequences. Alternatively, the mRNA sample is 10 separated by gel electrophoresis, transferred to a suitable support, e.g. nitrocellulose, nylon, etc., and then probed with a fragment of the subject DNA as a probe. Other techniques, such as oligonucleotide ligation assays, *in situ* hybridizations, and hybridization to DNA probes arrayed on a solid chip may also find use. Detection of mRNA hybridizing to the subject sequence is indicative of 15 Anthozoan protein gene expression in the sample.

The subject nucleic acids, including flanking promoter regions and coding regions, may be mutated in various ways known in the art to generate targeted. changes in promoter strength, sequence of the encoded protein, properties of the encoded protein, including fluorescent properties of the encoded protein, etc. The 20 DNA sequence or protein product of such a mutation will usually be substantially similar to the sequences provided herein, e.g. will differ by at least one nucleotide or amino acid, respectively, and may differ by at least two but not more than about ten nucleotides or amino acids. The sequence changes may be substitutions, insertions, deletions, or a combination thereof. Deletions may further include 25 larger changes, such as deletions of a domain or exon, e.g. of stretches of 10, 20, 50, 75, 100, 150 or more aa residues. Techniques for *in vitro* mutagenesis of cloned genes are known. Examples of protocols for site specific mutagenesis may be found in Gustin *et al.* (1993), *Biotechniques* 14:22; Barany (1985), *Gene* 37:111-23; Colicelli *et al.* (1985), *Mol. Gen. Genet.* 199:537-9; and Prentki *et al.* 30 (1984), *Gene* 29:303-13. Methods for site specific mutagenesis can be found in Sambrook *et al.*, *Molecular Cloning: A Laboratory Manual*, CSH Press 1989, pp. 15.3-15.108; Weiner *et al.* (1993), *Gene* 126:35-41; Sayers *et al.* (1992), *Biotechniques* 13:592-6; Jones and Winistorfer (1992), *Biotechniques* 12:528-30;

Barton *et al.* (1990), *Nucleic Acids Res* 18:7349-55; Marotti and Tomich (1989), *Gene Anal. Tech.* 6:67-70; and Zhu (1989), *Anal Biochem* 177:120-4. Such mutated nucleic acid derivatives may be used to study structure-function relationships of a particular chromo/ fluorescent protein, or to alter properties of the protein that affect its function or regulation.

5 Of particular interest in many embodiments is the following specific mutation protocol, which protocol finds use in mutating chromoproteins (e.g., colored proteins that have little if any fluorescence) into fluorescent mutants. In this protocol, the sequence of the candidate protein is aligned with the amino acid 10 sequence of *Aequorea victoria* wild type GFP, according to the protocol reported in Matz *et al.*, "Fluorescent proteins from non-bioluminescent Anthozoa species," *Nature Biotechnology* (October 1999) 17: 969 –973. Residue 148 of the aligned chromoprotein is identified and then changed to Ser, e.g., by site directed 15 mutagenesis, which results in the production of a fluorescent mutant of the wild type chromoprotein. See e.g., NFP-7 described below, which wild type protein is a chromoprotein that is mutated into a fluorescent protein by substitution of Ser for the native Ala residue at position 148.

Also of interest are humanized versions of the subject nucleic acids. As 20 used herein, the term "humanized" refers to changes made to the a nucleic acid sequence to optimize the codons for expression of the protein in human cells (Yang *et al.*, *Nucleic Acids Research* 24 (1996), 4592-4593). See also U.S. Patent No. 5,795,737 which describes humanization of proteins, the disclosure of which is herein incorporated by reference.

25 PROTEIN/POLYPEPTIDE COMPOSITIONS

Also provided by the subject invention are chromo- and/or fluorescent 30 proteins and mutants thereof, as well as polypeptide compositions related thereto. As the subject proteins are chromoproteins, they are colored proteins, which may be fluorescent, low or non- fluorescent. As used herein, the terms chromoprotein and fluorescent protein do not include luciferases, such as *Renilla luciferase*, and refer to any protein that is pigmented or colored and/or fluoresces when irradiated with light, e.g., white light or light of a specific wavelength (or narrow band of

wavelengths such as an excitation wavelength). The term polypeptide composition as used herein refers to both the full-length protein, as well as portions or fragments thereof. Also included in this term are variations of the naturally occurring protein, where such variations are homologous or substantially 5 similar to the naturally occurring protein, and mutants of the naturally occurring proteins, as described in greater detail below. The subject polypeptides are present in other than their natural environment.

In many embodiments, the subject proteins have an absorbance maximum ranging from about 300 to 700, usually from about 350 to 650 and more usually 10 from about 400 to 600 nm. Where the subject proteins are fluorescent proteins, by which is meant that they can be excited at one wavelength of light following which they will emit light at another wavelength, the excitation spectra of the subject proteins typically ranges from about 300 to 700, usually from about 350 to 650 and more usually from about 400 to 600 nm while the emission spectra of the subject 15 proteins typically ranges from about 400 to 800, usually from about 425 to 775 and more usually from about 450 to 750 nm. The subject proteins generally have a maximum extinction coefficient that ranges from about 10,000 to 50,000 and usually from about 15,000 to 45,000. The subject proteins typically range in length from about 150 to 300 and usually from about 200 to 300 amino acid residues; 20 and generally have a molecular weight ranging from about 15 to 35 kDa, usually from about 17.5 to 32.5 kDa.

In certain embodiments, the subject proteins are bright, where by bright is meant that the chromoproteins and their fluorescent mutants can be detected by common methods (e.g., visual screening, spectrophotometry, spectrofluorometry, 25 fluorescent microscopy, by FACS machines, etc.) Fluorescence brightness of particular fluorescent proteins is determined by its quantum yield multiplied by maximal extinction coefficient. Brightness of a chromoproteins may be expressed by its maximal extinction coefficient.

In certain embodiments, the subject proteins fold rapidly following 30 expression in the host cell. By rapidly folding is meant that the proteins achieve their tertiary structure that gives rise to their chromo- or fluorescent quality in a short period of time. In these embodiments, the proteins fold in a period of time that generally does not exceed about 3 days, usually does not exceed about 2 days and more usually does not exceed about 1 day.

Specific proteins of interest include the following specific proteins: (1) Green fluorescent protein from *Heteractis crispa* (hcriGFP); (2) Green fluorescent protein from *Dendronephthya* sp. (dendGFP); (3) Red fluorescent protein from *Zoanthus* sp. (zoanRFP); (4) Green fluorescent protein from *Scolymia cubensis* (scubGFP1);
5 (5) Green fluorescent protein from *Scolymia cubensis* (scubGFP2); (6) Red fluorescent protein from *Ricordea florida* (rfloRFP); (7) Green fluorescent protein from *Ricordea florida* (rfloGFP); (8) Red fluorescent protein from *Montastraea cavernosa* (mcavRFP); (9) Green fluorescent protein from *Montastraea cavernosa* (mcavGFP); (10) Green fluorescent protein from *Condylactis gigantea* (cgigGFP);
10 (11) Green fluorescent protein from *Agaricia fragilis* (afraGFP); (12) Green fluorescent protein from *Ricordea florida* (rfloGFP2); (13) Green fluorescent protein from *Montastraea cavernosa* (mcavGFP2); and (14) Green fluorescent protein homolog from *Montastraea annularis* (mannFP).

Homologs or proteins (or fragments thereof) that vary in sequence from the above provided specific amino acid sequences of the subject invention, i.e., SEQ ID NOS: 02; 04; 06; 08; 10; 12; 14; 16; 18; 20; 22; 24; 26 or 28, are also provided. By homolog is meant a protein having at least about 10%, usually at least about 20 % and more usually at least about 30 %, and in many embodiments at least about 35 %, usually at least about 40% and more usually at least about 60 % amino acid sequence identity to the protein of the subject invention, as determined using MegAlign, DNAsstar (1998) clustal algorithm as described in D. G. Higgins and P.M. Sharp, "Fast and Sensitive multiple Sequence Alignments on a Microcomputer," (1989) CABIOS, 5: 151-153. (Parameters used are ktuple 1, gap penalty 3, window, 5 and diagonals saved 5). In many embodiments, homologues of interest have much higher sequence identity, e.g., 65%, 70%, 75%, 80%, 85%, 90% or higher.

Also provided are proteins that are substantially identical to the wild type protein, where by substantially identical is meant that the protein has an amino acid sequence identity to the sequence of wild type protein of at least about 60%, usually at least about 65% and more usually at least about 70 %, where in some instances the identity may be much higher, e.g., 75%, 80%, 85%, 90%, 95% or higher.

In many embodiments, the subject homologues have structural features found in the above provided specific sequences, where such structural features include the β -can fold.

Proteins which are mutants of the above-described naturally occurring proteins are also provided. Mutants may retain biological properties of the wild-type (e.g., naturally occurring) proteins, or may have biological properties which differ from the wild-type proteins. The term "biological property" of the subject proteins includes, but is not limited to, spectral properties, such as absorbance maximum, emission maximum, maximum extinction coefficient, brightness (e.g., 5 as compared to the wild-type protein or another reference protein such as green fluorescent protein from *A. victoria*), and the like; *in vivo* and/or *in vitro* stability (e.g., half-life); etc. Mutants include single amino acid changes, deletions of one 10 or more amino acids, N-terminal truncations, C-terminal truncations, insertions, etc.

15 Mutants can be generated using standard techniques of molecular biology, e.g., random mutagenesis, and targeted mutagenesis. Several mutants are described herein. Given the guidance provided in the Examples, and using standard techniques, those skilled in the art can readily generate a wide variety of additional mutants and test whether a biological property has been altered. For 20 example, fluorescence intensity can be measured using a spectrophotometer at various excitation wavelengths.

Those proteins of the subject invention that are naturally occurring proteins are present in a non-naturally occurring environment, e.g., are separated from their naturally occurring environment. In certain embodiments, the subject proteins 25 are present in a composition that is enriched for the subject protein as compared to its naturally occurring environment. For example, purified protein is provided, where by purified is meant that the protein is present in a composition that is substantially free of non- chromo/fluoroprotein proteins of interest, where by substantially free is meant that less than 90 %, usually less than 60 % and more 30 usually less than 50 % of the composition is made up of non- chromoproteins or mutants thereof of interest. The proteins of the subject invention may also be present as an isolate, by which is meant that the protein is substantially free of other proteins and other naturally occurring biologic molecules, such as

oligosaccharides, polynucleotides and fragments thereof, and the like, where the term "substantially free" in this instance means that less than 70 %, usually less than 60% and more usually less than 50 % of the composition containing the isolated protein is some other naturally occurring biological molecule. In certain 5 embodiments, the proteins are present in substantially pure form, where by "substantially pure form" is meant at least 95%, usually at least 97% and more usually at least 99% pure.

In addition to the naturally occurring proteins, polypeptides that vary from the naturally occurring proteins, e.g., the mutant proteins described above, are 10 also provided. Generally such polypeptides include an amino acid sequence encoded by an open reading frame (ORF) of the gene encoding the subject wild type protein, including the full length protein and fragments thereof, particularly biologically active fragments and/or fragments corresponding to functional domains, and the like; and including fusions of the subject polypeptides to other 15 proteins or parts thereof. Fragments of interest will typically be at least about 10 aa in length, usually at least about 50 aa in length, and may be as long as 300 aa in length or longer, but will usually not exceed about 1000 aa in length, where the fragment will have a stretch of amino acids that is identical to the subject protein of at least about 10 aa, and usually at least about 15 aa, and in many embodiments 20 at least about 50 aa in length. In some embodiments, the subject polypeptides are about 25 aa, about 50 aa, about 75 aa, about 100 aa, about 125 aa, about 150 aa, about 200 aa, about 210 aa, about 220 aa, about 230 aa, or about 240 aa in length, up to the entire protein. In some embodiments, a protein fragment retains all or substantially all of a biological property of the wild-type protein.

25 The subject proteins and polypeptides may be obtained from naturally occurring sources or synthetically produced. For example, wild type proteins may be derived from biological sources which express the proteins, e.g., non-bioluminescent Cnidarian, e.g., Anthozoan, species, such as the specific ones listed above. The subject proteins may also be derived from synthetic means, e.g., 30 by expressing a recombinant gene or nucleic acid coding sequence encoding the protein of interest in a suitable host, as described above. Any convenient protein purification procedures may be employed, where suitable protein purification methodologies are described in Guide to Protein Purification, (Deuthser ed.) (Academic Press, 1990). For example, a lysate may be prepared from the original

source and purified using HPLC, exclusion chromatography, gel electrophoresis, affinity chromatography, and the like.

ANTIBODY COMPOSITIONS

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Also provided are antibodies that specifically bind to the subject fluorescent proteins. Suitable antibodies are obtained by immunizing a host animal with peptides comprising all or a portion of the subject protein. Suitable host animals include mouse, rat sheep, goat, hamster, rabbit, etc. The origin of the protein 10 immunogen will generally be a Cnidarian species, specifically a non-bioluminescent Cnidarian species, such as an Anthozoan species or a non-Petalucean Anthozoan species. The host animal will generally be a different species than the immunogen, e.g., mice, etc.

The immunogen may comprise the complete protein, or fragments and derivatives thereof. Preferred immunogens comprise all or a part of the protein, where these residues contain the post-translation modifications found on the native target protein. Immunogens are produced in a variety of ways known in the art; e.g., expression of cloned genes using conventional recombinant methods, isolation from Anthozoan species of origin, etc.

20 For preparation of polyclonal antibodies, the first step is immunization of the host animal with the target protein, where the target protein will preferably be in substantially pure form, comprising less than about 1% contaminant. The immunogen may comprise the complete target protein, fragments or derivatives thereof. To increase the immune response of the host animal, the target protein 25 may be combined with an adjuvant, where suitable adjuvants include alum, dextran, sulfate, large polymeric anions, oil & water emulsions, e.g. Freund's adjuvant, Freund's complete adjuvant, and the like. The target protein may also be conjugated to synthetic carrier proteins or synthetic antigens. A variety of hosts may be immunized to produce the polyclonal antibodies. Such hosts include rabbits, guinea pigs, rodents, e.g. mice, rats, sheep, goats, and the like. The target 30 protein is administered to the host, usually intradermally, with an initial dosage followed by one or more, usually at least two, additional booster dosages. Following immunization, the blood from the host will be collected, followed by

separation of the serum from the blood cells. The Ig present in the resultant antiserum may be further fractionated using known methods, such as ammonium salt fractionation, DEAE chromatography, and the like.

Monoclonal antibodies are produced by conventional techniques. Generally, 5 the spleen and/or lymph nodes of an immunized host animal provide a source of plasma cells. The plasma cells are immortalized by fusion with myeloma cells to produce hybridoma cells. Culture supernatant from individual hybridomas is screened using standard techniques to identify those producing antibodies with the desired specificity. Suitable animals for production of monoclonal antibodies to 10 the human protein include mouse, rat, hamster, etc. To raise antibodies against the mouse protein, the animal will generally be a hamster, guinea pig, rabbit, etc. The antibody may be purified from the hybridoma cell supernatants or ascites fluid by conventional techniques, e.g. affinity chromatography using protein bound to an insoluble support, protein A sepharose, etc.

15 The antibody may be produced as a single chain, instead of the normal multimeric structure. Single chain antibodies are described in Jost *et al.* (1994) J.B.C. 269:26267-73, and others. DNA sequences encoding the variable region of the heavy chain and the variable region of the light chain are ligated to a spacer encoding at least about 4 amino acids of small neutral amino acids, including 20 glycine and/or serine. The protein encoded by this fusion allows assembly of a functional variable region that retains the specificity and affinity of the original antibody.

Also of interest in certain embodiments are humanized antibodies. Methods of humanizing antibodies are known in the art. The humanized antibody 25 may be the product of an animal having transgenic human immunoglobulin constant region genes (see for example International Patent Applications WO 90/10077 and WO 90/04036). Alternatively, the antibody of interest may be engineered by recombinant DNA techniques to substitute the CH1, CH2, CH3, hinge domains, and/or the framework domain with the corresponding human 30 sequence (see WO 92/02190).

The use of Ig cDNA for construction of chimeric immunoglobulin genes is known in the art (Liu *et al.* (1987) P.N.A.S. 84:3439 and (1987) J. Immunol. 139:3521). mRNA is isolated from a hybridoma or other cell producing the

antibody and used to produce cDNA. The cDNA of interest may be amplified by the polymerase chain reaction using specific primers (U.S. Patent nos. 4,683,195 and 4,683,202). Alternatively, a library is made and screened to isolate the sequence of interest. The DNA sequence encoding the variable region of the antibody is then fused to human constant region sequences. The sequences of human constant regions genes may be found in Kabat *et al.* (1991) Sequences of Proteins of Immunological Interest, N.I.H. publication no. 91-3242. Human C region genes are readily available from known clones. The choice of isotype will be guided by the desired effector functions, such as complement fixation, or 5 activity in antibody-dependent cellular cytotoxicity. Preferred isotypes are IgG1, IgG3 and IgG4. Either of the human light chain constant regions, kappa or lambda, may be used. The chimeric, humanized antibody is then expressed by conventional methods.

10 Antibody fragments, such as Fv, F(ab')₂ and Fab may be prepared by cleavage of the intact protein, e.g. by protease or chemical cleavage. Alternatively, a truncated gene is designed. For example, a chimeric gene encoding a portion of the F(ab')₂ fragment would include DNA sequences encoding the CH1 domain and hinge region of the H chain, followed by a translational stop codon to yield the truncated molecule.

15 Consensus sequences of H and L J regions may be used to design oligonucleotides for use as primers to introduce useful restriction sites into the J region for subsequent linkage of V region segments to human C region segments. C region cDNA can be modified by site directed mutagenesis to place a restriction site at the analogous position in the human sequence.

20 Expression vectors include plasmids, retroviruses, YACs, EBV derived episomes, and the like. A convenient vector is one that encodes a functionally complete human CH or CL immunoglobulin sequence, with appropriate restriction sites engineered so that any VH or VL sequence can be easily inserted and expressed. In such vectors, splicing usually occurs between the splice donor site in the inserted J region and the splice acceptor site preceding the human C region, and also at the splice regions that occur within the human CH exons. 25 Polyadenylation and transcription termination occur at native chromosomal sites downstream of the coding regions. The resulting chimeric antibody may be joined 30

to any strong promoter, including retroviral LTRs, e.g. SV-40 early promoter, (Okayama *et al.* (1983) Mol. Cell. Bio. 3:280), Rous sarcoma virus LTR (Gorman *et al.* (1982) P.N.A.S. 79:6777), and moloney murine leukemia virus LTR (Grosschedl *et al.* (1985) Cell 41:885); native Ig promoters, etc.

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TRANSGENICS

The subject nucleic acids can be used to generate transgenic, non-human plants or animals or site specific gene modifications in cell lines. Transgenic cells of the subject invention include one or more nucleic acids according to the subject invention present as a transgene, where included within this definition are the parent cells transformed to include the transgene and the progeny thereof. In many embodiments, the transgenic cells are cells that do not normally harbor or contain a nucleic acid according to the subject invention. In those embodiments where the transgenic cells do naturally contain the subject nucleic acids, the nucleic acid will be present in the cell in a position other than its natural location, i.e. integrated into the genomic material of the cell at a non-natural location.

Transgenic animals may be made through homologous recombination, where the endogenous locus is altered. Alternatively, a nucleic acid construct is randomly integrated into the genome. Vectors for stable integration include plasmids, retroviruses and other animal viruses, YACs, and the like.

Transgenic organisms of the subject invention include cells and multicellular organisms, e.g., plants and animals, that are endogenous knockouts in which expression of the endogenous gene is at least reduced if not eliminated.

Transgenic organisms of interest also include cells and multicellular organisms, e.g., plants and animals, in which the protein or variants thereof is expressed in cells or tissues where it is not normally expressed and/or at levels not normally present in such cells or tissues.

DNA constructs for homologous recombination will comprise at least a portion of the gene of the subject invention, wherein the gene has the desired genetic modification(s), and includes regions of homology to the target locus. DNA constructs for random integration need not include regions of homology to mediate recombination. Conveniently, markers for positive and negative selection are included. Methods for generating cells having targeted gene modifications

through homologous recombination are known in the art. For various techniques for transfecting mammalian cells, see Keown *et al.* (1990), *Meth. Enzymol.* 185:527-537.

For embryonic stem (ES) cells, an ES cell line may be employed, or 5 embryonic cells may be obtained freshly from a host, e.g. mouse, rat, guinea pig, etc. Such cells are grown on an appropriate fibroblast-feeder layer or grown in the presence of leukemia inhibiting factor (LIF). When ES or embryonic cells have been transformed, they may be used to produce transgenic animals. After transformation, the cells are plated onto a feeder layer in an appropriate medium. 10 Cells containing the construct may be detected by employing a selective medium. After sufficient time for colonies to grow, they are picked and analyzed for the occurrence of homologous recombination or integration of the construct. Those colonies that are positive may then be used for embryo manipulation and blastocyst injection. Blastocysts are obtained from 4 to 6 week old superovulated 15 females. The ES cells are trypsinized, and the modified cells are injected into the blastocoel of the blastocyst. After injection, the blastocysts are returned to each uterine horn of pseudopregnant females. Females are then allowed to go to term and the resulting offspring screened for the construct. By providing for a different phenotype of the blastocyst and the genetically modified cells, chimeric progeny 20 can be readily detected.

The chimeric animals are screened for the presence of the modified gene and males and females having the modification are mated to produce homozygous progeny. If the gene alterations cause lethality at some point in development, tissues or organs can be maintained as allogeneic or congenic 25 grafts or transplants, or in *in vitro* culture. The transgenic animals may be any non-human mammal, such as laboratory animals, domestic animals, etc. The transgenic animals may be used in functional studies, drug screening, etc. Representative examples of the use of transgenic animals include those described infra.

30 Transgenic plants may be produced in a similar manner. Methods of preparing transgenic plant cells and plants are described in U.S. Pat. Nos. 5,767,367; 5,750,870; 5,739,409; 5,689,049; 5,689,045; 5,674,731; 5,656,466; 5,633,155; 5,629,470 ; 5,595,896; 5,576,198; 5,538,879; 5,484,956; the

disclosures of which are herein incorporated by reference. Methods of producing transgenic plants are also reviewed in Plant Biochemistry and Molecular Biology (eds Lea & Leegood, John Wiley & Sons)(1993) pp 275-295. In brief, a suitable plant cell or tissue is harvested, depending on the nature of the plant species. As 5 such, in certain instances, protoplasts will be isolated, where such protoplasts may be isolated from a variety of different plant tissues, e.g. leaf, hypocotyl, root, etc. For protoplast isolation, the harvested cells are incubated in the presence of cellulases in order to remove the cell wall, where the exact incubation conditions vary depending on the type of plant and/or tissue from which the cell is derived.

10 The resultant protoplasts are then separated from the resultant cellular debris by sieving and centrifugation. Instead of using protoplasts, embryogenic explants comprising somatic cells may be used for preparation of the transgenic host. Following cell or tissue harvesting, exogenous DNA of interest is introduced into the plant cells, where a variety of different techniques are available for such

15 introduction. With isolated protoplasts, the opportunity arise for introduction via DNA-mediated gene transfer protocols, including: incubation of the protoplasts with naked DNA, e.g. plasmids, comprising the exogenous coding sequence of interest in the presence of polyvalent cations, e.g.: PEG or PLO; and electroporation of the protoplasts in the presence of naked DNA comprising the exogenous sequence of interest. Protoplasts that have successfully taken up the exogenous DNA are then selected, grown into a callus, and ultimately into a 20 transgenic plant through contact with the appropriate amounts and ratios of stimulatory factors, e.g. auxins and cytokinins. With embryogenic explants, a convenient method of introducing the exogenous DNA in the target somatic cells is through the use of particle acceleration or "gene-gun" protocols. The resultant 25 explants are then allowed to grow into chimera plants, cross-bred and transgenic progeny are obtained. Instead of the naked DNA approaches described above, another convenient method of producing transgenic plants is *Agrobacterium* mediated transformation. With *Agrobacterium* mediated transformation, co- 30 integrative or binary vectors comprising the exogenous DNA are prepared and then introduced into an appropriate *Agrobacterium* strain, e.g. *A. tumefaciens*. The resultant bacteria are then incubated with prepared protoplasts or tissue explants, e.g. leaf disks, and a callus is produced. The callus is then grown under selective

conditions, selected and subjected to growth media to induce root and shoot growth to ultimately produce a transgenic plant.

UTILITY

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The subject chromoproteins and fluorescent mutants thereof find use in a variety of different applications, where the applications necessarily differ depending on whether the protein is a chromoprotein or a fluorescent protein.

Representative uses for each of these types of proteins will be described below,

10 where the follow described uses are merely representative and are in no way meant to limit the use of the subject proteins to those described below.

Chromoproteins

15 The subject chromoproteins of the present invention find use in a variety of different applications. One application of interest is the use of the subject proteins as coloring agents which are capable of imparting color or pigment to a particular composition of matter. Of particular interest in certain embodiments are non-toxic chromoproteins. The subject chromoproteins may be incorporated into a variety of 20 different compositions of matter, where representative compositions of matter include: food compositions, pharmaceuticals, cosmetics, living organisms, e.g., animals and plants, and the like. Where used as a coloring agent or pigment, a sufficient amount of the chromoprotein is incorporated into the composition of matter to impart the desired color or pigment thereto. The chromoprotein may be 25 incorporated into the composition of matter using any convenient protocol, where the particular protocol employed will necessarily depend, at least in part, on the nature of the composition of matter to be colored. Protocols that may be employed include, but are not limited to: blending, diffusion, friction, spraying, injection, tattooing, and the like.

30 The chromoproteins may also find use as labels in analyte detection assays, e.g., assays for biological analytes of interest. For example, the chromoproteins may be incorporated into adducts with analyte specific antibodies or binding fragments thereof and subsequently employed in immunoassays for

analytes of interest in a complex sample, as described in U.S. Patent No. 4,302,536; the disclosure of which is herein incorporated by reference. Instead of antibodies or binding fragments thereof, the subject chromoproteins or chromogenic fragments thereof may be conjugated to ligands that specifically bind to an analyte of interest, or other moieties, growth factors, hormones, and the like; as is readily apparent to those of skill in the art.

In yet other embodiments, the subject chromoproteins may be used as selectable markers in recombinant DNA applications, e.g., the production of transgenic cells and organisms, as described above. As such, one can engineer a particular transgenic production protocol to employ expression of the subject chromoproteins as a selectable marker, either for a successful or unsuccessful protocol. Thus, appearance of the color of the subject chromoprotein in the phenotype of the transgenic organism produced by a particular process can be used to indicate that the particular organism successfully harbors the transgene of interest, often integrated in a manner that provides for expression of the transgene in the organism. When used a selectable marker, a nucleic acid encoding for the subject chromoprotein can be employed in the transgenic generation process, where this process is described in greater detail supra. Particular transgenic organisms of interest where the subject proteins may be employed as selectable markers include transgenic plants, animals, bacteria, fungi, and the like.

In yet other embodiments, the chromoproteins (and fluorescent proteins) of the subject invention find use in sunscreens, as selective filters, etc., in a manner similar to the uses of the proteins described in WO 00/46233.

25 *Fluorescent Proteins*

The subject fluorescent proteins of the present invention (as well as other components of the subject invention described above) find use in a variety of different applications, where such applications include, but are not limited to, the following. The first application of interest is the use of the subject proteins in fluorescence resonance energy transfer (FRET) applications. In these applications, the subject proteins serve as donor and/or acceptors in combination with a second fluorescent protein or dye, e.g., a fluorescent protein as described in

Matz et al., *Nature Biotechnology* (October 1999) 17:969-973, a green fluorescent protein from *Aequoria victoria* or fluorescent mutant thereof, e.g., as described in U.S. Patent No. 6,066,476; 6,020,192; 5,985,577; 5,976,796; 5,968,750; 5,968,738; 5,958,713; 5,919,445; 5,874,304, the disclosures of which are herein incorporated by reference, other fluorescent dyes, e.g., coumarin and its derivatives, e.g. 7-amino-4-methylcoumarin, aminocoumarin, bodipy dyes, such as Bodipy FL, cascade blue, fluorescein and its derivatives, e.g. fluorescein isothiocyanate, Oregon green, rhodamine dyes, e.g. texas red, tetramethylrhodamine, eosins and erythrosins, cyanine dyes, e.g. Cy3 and Cy5, 10 macrocyclic chelates of lanthanide ions, e.g. quantum dye, etc., chemiluminescent dyes, e.g., luciferases, including those described in U.S. Patent Nos. 5,843,746; 5,700,673; 5,674,713; 5,618,722; 5,418,155; 5,330,906; 5,229,285; 5,221,623; 5,182,202; the disclosures of which are herein incorporated by reference. Specific examples of where FRET assays employing the subject fluorescent proteins may be used include, but are not limited to: the detection of protein-protein interactions, e.g., mammalian two-hybrid system, transcription factor dimerization, membrane protein multimerization; multiprotein complex formation, etc., as a biosensor for a number of different events, where a peptide or protein covalently links a FRET fluorescent combination including the subject fluorescent proteins and the linking peptide or protein is, e.g., a protease specific substrate, e.g., for caspase 20 mediated cleavage, a linker that undergoes conformational change upon receiving a signal which increases or decreases FRET, e.g., PKA regulatory domain (cAMP-sensor), phosphorylation, e.g., where there is a phosphorylation site in the linker or the linker has binding specificity to phosphorylated/dephosphorylated domain of another protein, or the linker has Ca^{2+} binding domain. Representative fluorescence resonance energy transfer or FRET applications in which the subject proteins find use include, but are not limited to, those described in: U.S. Patent Nos. 6,008,373; 5,998,146; 5,981,200; 5,945,526; 5,945,283; 5,911,952; 5,869,255; 5,866,336; 5,863,727; 5,728,528; 5,707,804; 5,688,648; 5,439,797; the disclosures of which are herein incorporated by reference.

The subject fluorescent proteins also find use as biosensors in prokaryotic and eukaryotic cells, e.g. as Ca^{2+} ion indicator; as pH indicator, as phosphorylation indicator, as an indicator of other ions, e.g., magnesium, sodium, potassium, chloride and halides. For example, for detection of Ca ion, proteins containing an

EF-hand motif are known to translocate from the cytosol to membranes upon Ca^{2+} binding. These proteins contain a myristoyl group that is buried within the molecule by hydrophobic interactions with other regions of the protein. Binding of Ca^{2+} induces a conformational change exposing the myristoyl group which then is 5 available for the insertion into the lipid bilayer (called a " Ca^{2+} -myristoyl switch"). Fusion of such a EF-hand containing protein to Fluorescent Proteins (FP) could make it an indicator of intracellular Ca^{2+} by monitoring the translocation from the cytosol to the plasma membrane by confocal microscopy. EF-hand proteins suitable for use in this system include, but are not limited to: recoverin (1-3), 10 calcineurin B, troponin C, visinin, neurocalcin, calmodulin, parvalbumin, and the like. For pH, a system based on hisactophilins may be employed. Hisactophilins are myristoylated histidine-rich proteins known to exist in *Dictyostelium*. Their binding to actin and acidic lipids is sharply pH-dependent within the range of cytoplasmic pH variations. In living cells membrane binding seems to override the 15 interaction of hisactophilins with actin filaments. At $\text{pH} \leq 6.5$ they locate to the plasma membrane and nucleus. In contrast, at $\text{pH} 7.5$ they evenly distribute throughout the cytoplasmic space. This change of distribution is reversible and is attributed to histidine clusters exposed in loops on the surface of the molecule. The reversion of intracellular distribution in the range of cytoplasmic pH variations 20 is in accord with a pK of 6.5 of histidine residues. The cellular distribution is independent of myristylation of the protein. By fusing FPs (Fluorescent Proteins) to hisactophilin the intracellular distribution of the fusion protein can be followed by laser scanning, confocal microscopy or standard fluorescence microscopy. Quantitative fluorescence analysis can be done by performing line scans through 25 cells (laser scanning confocal microscopy) or other electronic data analysis (e.g., using metamorph software (Universal Imaging Corp) and averaging of data collected in a population of cells. Substantial pH-dependent redistribution of hisactophilin-FP from the cytosol to the plasma membrane occurs within 1-2 min and reaches a steady state level after 5-10 min. The reverse reaction takes place 30 on a similar time scale. As such, hisactophilin-fluorescent protein fusion protein that acts in an analogous fashion can be used to monitor cytosolic pH changes in real time in live mammalian cells. Such methods have use in high throughput applications, e.g., in the measurement of pH changes as consequence of growth factor receptor activation (e.g. epithelial or platelet-derived growth factor)

chemotactic stimulation/ cell locomotion, in the detection of intracellular pH changes as second messenger, in the monitoring of intracellular pH in pH manipulating experiments, and the like. For detection of PKC activity, the reporter system exploits the fact that a molecule called MARCKS (myristoylated alanine-rich C kinase substrate) is a PKC substrate. It is anchored to the plasma membrane via myristylation and a stretch of positively charged amino acids (ED-domain) that bind to the negatively charged plasma membrane via electrostatic interactions. Upon PKC activation the ED-domain becomes phosphorylated by PKC, thereby becoming negatively charged, and as a consequence of electrostatic repulsion MARCKS translocates from the plasma membrane to the cytoplasm (called the "myristoyl-electrostatic switch"). Fusion of the N-terminus of MARCKS ranging from the myristylation motif to the ED-domain of MARCKS to fluorescent proteins of the present invention makes the above a detector system for PKC activity. When phosphorylated by PKC, the fusion protein translocates from the plasma membrane to the cytosol. This translocation is followed by standard fluorescence microscopy or confocal microscopy e.g. using the Cellomics technology or other High Content Screening systems (e.g. Universal Imaging Corp./Becton Dickinson). The above reporter system has application in High Content Screening, e.g., screening for PKC inhibitors, and as an indicator for PKC activity in many screening scenarios for potential reagents interfering with this signal transduction pathway. Methods of using fluorescent proteins as biosensors also include those described in U.S. Patent Nos. 972,638; 5,824,485 and 5,650,135 (as well as the references cited therein) the disclosures of which are herein incorporated by reference.

The subject fluorescent proteins also find use in applications involving the automated screening of arrays of cells expressing fluorescent reporting groups by using microscopic imaging and electronic analysis. Screening can be used for drug discovery and in the field of functional genomics: e.g., where the subject proteins are used as markers of whole cells to detect changes in multicellular reorganization and migration, e.g., formation of multicellular tubules (blood vessel formation) by endothelial cells, migration of cells through Fluoroblok Insert System (Becton Dickinson Co.), wound healing, neurite outgrowth, etc.; where the proteins are used as markers fused to peptides (e.g., targeting sequences) and proteins that allow the detection of change of intracellular location as indicator for cellular

activity, for example: signal transduction, such as kinase and transcription factor translocation upon stimuli, such as protein kinase C, protein kinase A, transcription factor NFkB, and NFAT; cell cycle proteins, such as cyclin A, cyclin B1 and cyclinE; protease cleavage with subsequent movement of cleaved substrate, phospholipids, with markers for intracellular structures such as endoplasmic reticulum, Golgi apparatus, mitochondria, peroxisomes, nucleus, nucleoli, plasma membrane, histones, endosomes, lysosomes, microtubules, actin) as tools for High Content Screening: co-localization of other fluorescent fusion proteins with these localization markers as indicators of movements of intracellular fluorescent fusion proteins/peptides or as marker alone; and the like. Examples of applications involving the automated screening of arrays of cells in which the subject fluorescent proteins find use include: U.S. Patent No. 5,989,835; as well as WO/0017624; WO 00/26408; WO 00/17643; and WO 00/03246; the disclosures of which are herein incorporated by reference.

The subject fluorescent proteins also find use in high through-put screening assays. The subject fluorescent proteins are stable proteins with half-lives of more than 24h. Also provided are destabilized versions of the subject fluorescent proteins with shorter half-lives that can be used as transcription reporters for drug discovery. For example, a protein according to the subject invention can be fused with a putative proteolytic signal sequence derived from a protein with shorter half-life, e.g., PEST sequence from the mouse ornithine decarboxylase gene, mouse cyclin B1 destruction box and ubiquitin, etc. For a description of destabilized proteins and vectors that can be employed to produce the same, see e.g., U.S. Patent No. 6,130,313; the disclosure of which is herein incorporated by reference.

Promoters in signal transduction pathways can be detected using destabilized versions of the subject fluorescent proteins for drug screening, e.g., AP1, NFAT, NFkB, Smad, STAT, p53, E2F, Rb, myc, CRE, ER, GR and TRE, and the like.

The subject proteins can be used as second messenger detectors, e.g., by fusing the subject proteins to specific domains: e.g., PKCgamma Ca binding domain, PKCgamma DAG binding domain, SH2 domain and SH3 domain, etc.

Secreted forms of the subject proteins can be prepared, e.g. by fusing secreted leading sequences to the subject proteins to construct secreted forms of the subject proteins, which in turn can be used in a variety of different applications.

The subject proteins also find use in fluorescence activated cell sorting applications. In such applications, the subject fluorescent protein is used as a label to mark a population of cells and the resulting labeled population of cells is then sorted with a fluorescent activated cell sorting device, as is known in the art. FACS methods are described in U.S. Patent Nos. 5,968,738 and 5,804,387; the disclosures of which are herein incorporated by reference.

The subject proteins also find use as *in vivo* marker in animals (e.g., transgenic animals). For example, expression of the subject protein can be driven by tissue specific promoters, where such methods find use in research for gene 10 therapy, e.g., testing efficiency of transgenic expression, among other applications. A representative application of fluorescent proteins in transgenic animals that illustrates this class of applications of the subject proteins is found in WO 00/02997, the disclosure of which is herein incorporated by reference.

Additional applications of the subject proteins include: as markers following 15 injection into cells or animals and in calibration for quantitative measurements (fluorescence and protein); as markers or reporters in oxygen biosensor devices for monitoring cell viability; as markers or labels for animals, pets, toys, food, etc.; and the like.

The subject fluorescent proteins also find use in protease cleavage assays. 20 For example, cleavage inactivated fluorescence assays can be developed using the subject proteins, where the subject proteins are engineered to include a protease specific cleavage sequence without destroying the fluorescent character of the protein. Upon cleavage of the fluorescent protein by an activated protease fluorescence would sharply decrease due to the destruction of a functional 25 chromophor. Alternatively, cleavage activated fluorescence can be developed using the subject proteins, where the subject proteins are engineered to contain an additional spacer sequence in close proximity/or inside the chromophor. This variant would be significantly decreased in its fluorescent activity, because parts of the functional chromophor would be divided by the spacer. The spacer would be 30 framed by two identical protease specific cleavage sites. Upon cleavage via the activated protease the spacer would be cut out and the two residual "subunits" of the fluorescent protein would be able to reassemble to generate a functional fluorescent protein. Both of the above types of application could be developed in assays for a variety of different types of proteases, e.g., caspases, etc.

The subject proteins can also be used in assays to determine the phospholipid composition in biological membranes. For example, fusion proteins of the subject proteins (or any other kind of covalent or non-covalent modification of the subject proteins) that allows binding to specific phospholipids to 5 localize/visualize patterns of phospholipid distribution in biological membranes also allowing colocalization of membrane proteins in specific phospholipid rafts can be accomplished with the subject proteins. For example, the PH domain of GRP1 has a high affinity to phosphatidyl-inositol tri-phosphate (PIP3) but not to PIP2. As such, a fusion protein between the PH domain of GRP1 and the subject 10 proteins can be constructed to specifically label PIP3 rich areas in biological membranes.

Yet another application of the subject proteins is as a fluorescent timer, in which the switch of one fluorescent color to another (e.g. green to red) concomitant with the ageing of the fluorescent protein is used to determine the 15 activation/deactivation of gene expression, e.g., developmental gene expression, cell cycle dependent gene expression, circadian rhythm specific gene expression, and the like.

The antibodies of the subject invention, described above, also find use in a number of applications, including the differentiation of the subject proteins from 20 other fluorescent proteins.

KITS

Also provided by the subject invention are kits for use in practicing one or 25 more of the above described applications, where the subject kits typically include elements for making the subject proteins, e.g., a construct comprising a vector that includes a coding region for the subject protein. The subject kit components are typically present in a suitable storage medium, e.g., buffered solution, typically in a suitable container. Also present in the subject kits may be antibodies to the 30 provided protein. In certain embodiments, the kit comprises a plurality of different vectors each encoding the subject protein, where the vectors are designed for expression in different environments and/or under different conditions, e.g., constitutive expression where the vector includes a strong promoter for expression

in mammalian cells, a promoterless vector with a multiple cloning site for custom insertion of a promoter and tailored expression, etc.

In addition to the above components, the subject kits will further include instructions for practicing the subject methods. These instructions may be present in the subject kits in a variety of forms, one or more of which may be present in the kit. One form in which these instructions may be present is as printed information on a suitable medium or substrate, e.g., a piece or pieces of paper on which the information is printed, in the packaging of the kit, in a package insert, etc. Yet another means would be a computer readable medium, e.g., diskette, CD, etc., on which the information has been recorded. Yet another means that may be present is a website address which may be used via the internet to access the information at a removed site. Any convenient means may be present in the kits.

15

The following examples are offered by way of illustration and not by way of limitation.

EXPERIMENTAL

25 I. Introduction

In the following experimental section, we present eleven new GFP-like proteins.

30 II. Materials and Methods

A. Collection of samples

Samples (100-500 mg of tissue) of *Montastraea cavernosa*, *Condylactis gigantea*, *Scolymia cubensis* and *Ricordea florida* were collected at Florida Keys

Marine Sanctuary (Long Key), under National Marine Sanctuary authorization FKNMS-2000-009. The samples were collected during night dives, candidate specimens were picked on the basis of their appearance under ultraviolet flashlight. Other samples (*Dendronephthya* sp., *Heteractis crispa*, *Discosoma* sp.3, 5 *Zoanthus* sp. 2) were picked from private seawater aquariums.

B. Cloning and expression of GFP-like proteins

Total RNA was isolated from the tissue samples following the protocol described in Chomczynski, P. & Sacchi, N. (1987) *Anal Biochem* **162**, 156-9. Total 10 cDNA was amplified using SMART™ cDNA amplification kit (Clontech). These amplified cDNA samples were used to amplify 3'-fragments of cDNAs coding for GFP-like proteins and then obtain the missing 5'-flanks, exactly as described in Matz, M. V., Fradkov, A. F., Labas, Y. A., Savitsky, A. P., Zaraisky, A. G., Markelov, M. L. & Lukyanov, S. A. (1999) *Nat Biotechnol* **17**, 969-73.. After 15 determining the complete cDNA sequence, the coding regions were amplified using the same cDNA samples as were used to clone the 3'- and 5'-flanks as templates. An upstream ("N-terminal") primer had a 5'-heel (5'-tTGAtTGAtTGAAGGAGAatatc) carrying stop codons (bold) in all frames and bacterial ribosome-binding site (underlined), followed by the target cDNA 20 sequence (20-22 bases) starting with initiation codon of the ORF. The downstream ("C-terminal") primer was 22-25 bases long and corresponded to the antisense sequence of cDNA around the stop codon of the ORF. The resulting fragments were cloned using pGEM-T vector cloning kit (Promega) following the manufacturer's protocol, using *Escherichia coli* JM109 strain as host. The colonies 25 were grown on LB/agar/carbenicillin plates supplemented with 0.3 mM IPTG for 16-20 hours at 37°C, and then incubated for two days at 4°C. The fluorescent colonies were selected using fluorescent microscope and streaked widely on new plates. The same colonies were used for overnight culture inoculation followed by plasmid isolation and sequencing, to confirm the identity of the clone. The bacteria 30 were harvested from the plates, suspended in 1 ml of PBS and disrupted by sonication. The lysate was cleared by centrifugation, and its fluorescent properties were determined using LS-50B spectrofluorometer (Perkin Elmer Instruments). For

mcavRFP and rfloGFP, the "early" samples were harvested after 24 hours at 37°C, "late" samples – after 24 hours at 37C followed by four days at 4°C.

C. Phylogenetic analysis

5 The alignment of GFP-like proteins (see supplemental data) was constructed after Matz, M. V., Fradkov, A. F., Labas, Y. A., Savitsky, A. P., Zaraisky, A. G., Markelov, M. L. & Lukyanov, S. A. (1999) *Nat Biotechnol* **17**, 969-73 taking in account constraints of the protein structure. Then the DNA alignment was made following the protein alignment; excluding the poorly aligned N- and C-10 terminal regions. The phylogenetic tree was constructed using Tree-Puzzle software (Strimmer, K. & von Haeseler, A. (1996) *Mol. Biol. Evol.* **13**, 964-969) under HKY model of DNA evolution (Hasegawa, M., Kishino, H. & Yano, K. (1985) *J. Mol. Evol.* **22**, 160-174), assuming that the variability of sites follows gamma-distribution with alpha parameter estimated from the dataset. The tree was 15 confirmed to be the maximum likelihood tree by PAML software (Yang, Z. (2000) (University College (<http://abacus.gene.ucl.ac.uk/software/paml.html>), London, England) under REV model (Yang, Z. H., Goldman, N. & Friday, A. (1994) *Molecular Biology and Evolution* **11**, 316-324). The tree built by Tree-Puzzle from protein alignment (JTT model, (Jones, D. T., Taylor, W. R. & Thornton, J. M. 20 (1992) *CABIOS* **8**, 275-282) had the same topology but lower support values due to smaller number of informative sites in the protein alignment.

III. Results and Discussion

25 A. Nomenclature

For the sake of clarity of phylogenetic analysis representation, in this paper we are using new nomenclature for GFP-like proteins. Our protein identification tags include four-letter leader composed of first letter of genus name and three initial letters of species name, followed by definition of color type: GFP – green, 30 RFP – red, YFP – yellow, CP – chromoprotein (non-fluorescent). When the species is not defined, the leader is four initial letters of the genus name. In the case of multiple non-identified species of the same genus, a number is added to the leader (such as in dis3GFP or zoan2RFP); in the case of several proteins of

the same color type found in the same species, the number is added to the color definition (such as in scubGFP1 and scubGFP2). For *Aequorea victoria* GFP and drFP583 from *Discosoma* sp., widely accepted common names are kept: GFP and DsRed.

5

B. New GFP-like proteins

A total of fourteen new GFP-like proteins were cloned and spectroscopically characterized. The spectral features of 11 of these proteins are summarized in Table 1 appearing in the figures, as well as the other figures of the application

10 This subset of 11 includes representatives exhibiting features not seen before in Anthozoan GFP-like proteins. Two green proteins from *Condylactis gigantea* (cgigGFP) and *Heteractis crispa* (hcriGFP) possess double-peaked excitation spectra very similar to the one of wild-type GFP, suggesting that their chromophores undergo photoconversion between neutral and ionized states (

15 Brejc, K., Sixma, T. K., Kitts, P. A., Kain, S. R., Tsien, R. Y., Ormo, M. & Remington, S. J. (1997) *Proc Natl Acad Sci U S A* **94**, 2306-11; Palm, G. J., Zdanov, A., Gaitanaris, G. A., Stauber, R., Pavlakis, G. N. & Wlodawer, A. (1997) *Nat Struct Biol* **4**, 361-5). The red-emitting protein zoan2RFP, although being very similar to DsRed in the shape of excitation/emission curves, behaves like "timer": it turns green at first and then matures into red (Fig. 1, A and B), similarly to one of the mutant variants of DsRed (Terskikh, A., Fradkov, A., Ermakova, G., Zaraisky, A., Tan, P., Kajava, A. V., Zhao, X., Lukyanov, S., Matz, M., Kim, S., Weissman, I. & Siebert, P. (2000) *Science* **290**, 1585-8.). The two new red-emitters from great star coral *Montastraea cavernosa* (mcavRFP) and florida corallimorph *Ricordea florida* (rfloRFP) also show a "timer" phenotype (Fig. 1, C-F). In contrast to zoan2RFP, they failed to mature completely into red in our bacterial expression trials, which resulted in two-peak emission spectra such as shown in Figure 1 (D and F). Remarkably, for both these proteins, the red emission band in the more mature form had major excitation peak virtually identical to the one of the immature green form, the yellow-orange excitation peak being significantly smaller (Fig. 2). This is strikingly different from the rest of the orange-red proteins, in which the red emission is excited best in yellow-orange region (Figure 4, Table 1, spectra E). This unusual shape of excitation spectra may be due to photoconversion of the

ionization states of the chromophore (by analogy with green proteins), or to even more profound differences in the chromophore structure. In favor of the latter speaks the fact that the shape of the red emission peaks of mcavRFP and rfloRFP is notably different from other orange-red proteins: it is much narrower and almost symmetrical in contrast to the wide and skewed emission peak of the others (compare spectra E and F in Table 1, Figure 4). Meanwhile, in GFP from *Aequorea victoria*, presence or absence of photoconversion does not have much effect on the shape of emission spectra (Heim, R., Cubitt, A. B. & Tsien, R. Y. (1995) *Nature* **373**, 663-4). The striking similarity of major excitation peaks for mature and immature proteins makes it tempting to suggest that in mcavRFP and rfloRFP, the "built-in" fluorescence resonance energy transfer (FRET) from immature green form of the protein to the mature red form is the major mechanism giving rise to red emission.

15 C. Structural/spectral types of GFP-like proteins

In our view, the best way to classify GFP-like proteins is by their color as it appears to human eye. We discriminate four color types of GFP-like proteins: green, yellow, orange-red and purple-blue, or chromoproteins (Table 1, Figure 14). All of them share the same fold of polypeptide chain, termed "beta-can" (Ormo, M., Cubitt, A. B., Kallio, K., Gross, L. A., Tsien, R. Y. & Remington, S. J. (1996) *Science* **273**, 1392-5.; Yang, F., Moss, L. G. & Phillips, G. N., Jr. (1996) *Nat Biotechnol* **14**, 1246-51). However, there are substantial differences between these color types as far as the chromophore structure is concerned (see Table 1). In GFP (green color), the chromophore is formed by residues 65-67 (Ser-Tyr-Gly) as a result of condensation between the carbonyl carbon of Ser-65 and the amino nitrogen of Gly-67 that produces a five-member ring, followed by the dehydrogenation of the Tyr-66 methylene bridge. All the green proteins apparently possess the same chromophore, and the differences in the spectral shapes are explained by modifications of its environment. It must be noted that the green proteins having excitation/emission spectra such as on panel A on Table 1 are sometimes called cyan or even blue, but to the human eye the color of these proteins after purification still appears bright green. In the red protein DsRed, the chromophore synthesis includes one more stage that extends the conjugated pi-

system of the chromophore – dehydrogenation of the bond between the alpha carbon and amino nitrogen of the first chromophore-forming residue. Meanwhile, in the chromoproteins representative asulCP, cyclization leads to the formation of a six-member rather than five-member ring, and the critical step in creating the 5 extended conjugated pi-system is breakage of the polypeptide chain immediately before the chromophore. Notably, no other chromoprotein contains such a chain break, as demonstrated by denaturing electrophoresis of the bacterial exprssion products (data not shown). This indicates that the chromophore structure of asulCP is exception rather than the rule within this color type. Biochemical and 10 mutagenesis studies of the yellow zoanYFP indicated that this protein has yet another chromophore structure. So, it must be concluded that although pronounced color difference between GFP-like proteins indicates difference in chromophore structures (which makes it reasonable to use color for classification), different chromophores might be found even in the proteins of the same color, as 15 it happens within the group of chromoproteins and probably within the orange-red group.

D. Molecular basis of color conversion

Since a chromophore synthesis pathway in DsRed is an extended form of 20 the GFP pathway, it can be easily imagined that any mutation damaging the additional autocatalytic stage in DsRed would convert it into green protein. Indeed, at least seven different mutant variants of DsRed emitting in the green range were found during random and site specific mutagenesis. Similar reasoning should apply to the two new red proteins, because their red emission also arises as a 25 result of further modification of the green-emitting chromophore.

It has been shown that a single amino acid replacement can convert a chromoprotein into a DsRed-like red fluorescent protein. It is particularly unexpected for asulCP from *Anemonia sulcata*, which has been directly demonstrated to contain a very dissimilar chromophore; and it still seems unlikely 30 that its red fluorescent mutant variant actually switches to synthesizing a DsRed-type chromophore instead of original one. However, random mutations in this mutant variant resulted in appearance of green-emitting forms. Since no green-emitting intermediate stage was present in the original asulCP autocatalytic pathway, formation of green-emitting structure in these mutants signifies a

substantial deviation, most probably towards a GFP/DsRed type of chromophore formation sequence judging by the shape of excitation/emission spectra of the green asulCP mutants.

5 Finally, yellow protein zoanYFP also can be converted into green-emitting state by at least two different amino acid replacements.

Taking these data into account, the following explanation of the observed phylogenetic pattern seems plausible: that different chromophore structures, even the most dissimilar ones, are alternative products synthesized with the help of a basically similar autocatalytic environment, rather than outcomes of prolonged 10 evolution of different catalytic mechanisms. Apparently, just a few amino acid changes in the protein may act like a switch between alternative pathways, as exemplified by mutagenesis results on asulCP chromoprotein.

All publications and patent applications cited in this specification are herein 15 incorporated by reference as if each individual publication or patent application were specifically and individually indicated to be incorporated by reference. The citation of any publication is for its disclosure prior to the filing date and should not be construed as an admission that the present invention is not entitled to antedate such publication by virtue of prior invention.

20 Although the foregoing invention has been described in some detail by way of illustration and example for purposes of clarity of understanding, it is readily apparent to those of ordinary skill in the art in light of the teachings of this invention that certain changes and modifications may be made thereto without departing from the spirit or scope of the appended claims.

WHAT IS CLAIMED IS:

1. A nucleic acid having a sequence of residues that is substantially the same as or identical to a nucleotide sequence of at least 10 residues in length of SEQ ID NOS:01, 03, 05, 07, 09, 11, 13, 15, 17, 19, 21, 23, 25 or 27.

5

2. The nucleic acid according to Claim 1, wherein said nucleic acid has a sequence similarity of at least about 60% with a sequence of at least 10 residues in length of SEQ ID NOS: 01, 03, 05, 07, 09, 11, 13, 15, 17, 19, 21, 23, 25 or 27.

10

3. A nucleic acid present in other than its natural environment that encodes a chromo and/or fluorescent protein that has an amino acid sequence of: SEQ ID NOS: 02, 04, 06, 08, 10, 12, 14, 16, 18, 20, 22, 24, 26 or 28.

15

4. A nucleic acid that encodes a mutant protein of a protein that has an amino acid sequence of: SEQ ID NOS: 02, 04, 06, 08, 10, 12, 14, 16, 18, 20, 22, 24, 26 or 28.

5. The nucleic acid according to Claim 4, wherein said mutant protein comprises at least one point mutation as compared to its wild type protein.

20

6. The nucleic acid according to Claim 4, wherein said mutant protein comprises at least one deletion mutation as compared to its wild type protein.

7. A fragment of the nucleic acid selected of Claims 1 to 6.

25

8. An isolated nucleic acid or mimetic thereof that hybridizes under stringent conditions to a nucleic acid of Claims 1 to 7.

9. A construct comprising a vector and a nucleic acid of Claims 1 to 8.

30

10. An expression cassette comprising:

- (a) a transcriptional initiation region functional in an expression host;
- (b) a nucleic acid selected from the group consisting of the nucleic acids of Claims 1 to 9; and

(c) and a transcriptional termination region functional in said expression host.

11. A cell, or the progeny thereof, comprising an expression cassette according to Claim 10 as part of an extrachromosomal element or integrated into the genome of a host cell as a result of introduction of said expression cassette into said host cell.

12. A method of producing a chromo and/or fluorescent protein, said method comprising:

growing a cell according to Claim 11, whereby said protein is expressed; and

isolating said protein substantially free of other proteins.

13. A protein or fragment thereof encoded by a nucleic acid selected from the group consisting of Claims 1 to 10.

14. An antibody binding specifically to a protein according to Claim 13.

15. A transgenic cell or the progeny thereof comprising a transgene selected from the group consisting of a nucleic acids according to any of Claims 1 to 10.

16. A transgenic organism capable comprising a transgene selected from the group consisting of a nucleic acids according to any of Claims 1 to 10.

17. In an application that employs a chromo- or fluorescent protein, the improvement comprising:

employing a protein according to Claim 13.

18. In an application that employs a nucleic acid encoding a chromo- or fluorescent protein, the improvement comprising:

employing a nucleic acid according to Claims 1 to 10.

19. A kit comprising a nucleic acid according to Claims 1 to 10 and instructions for using said nucleic acid.

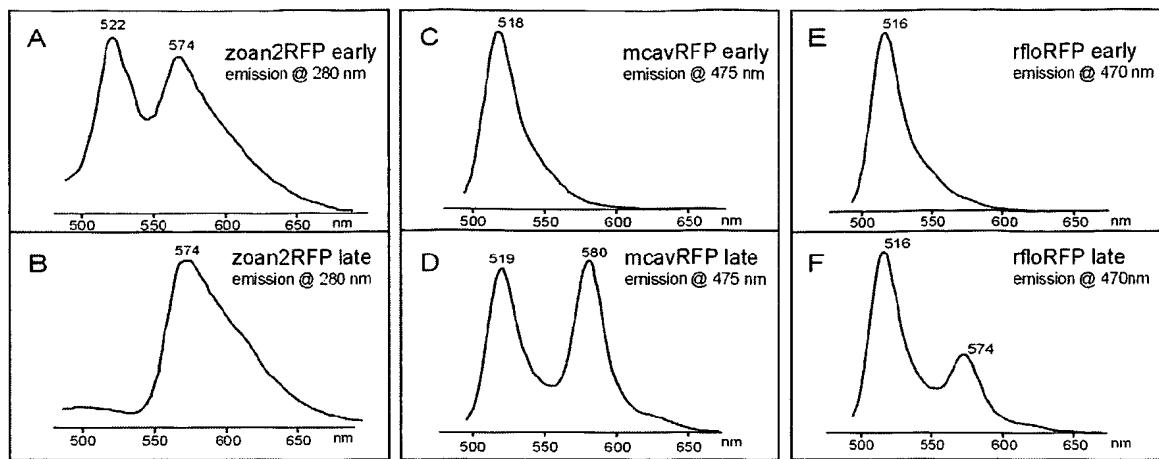
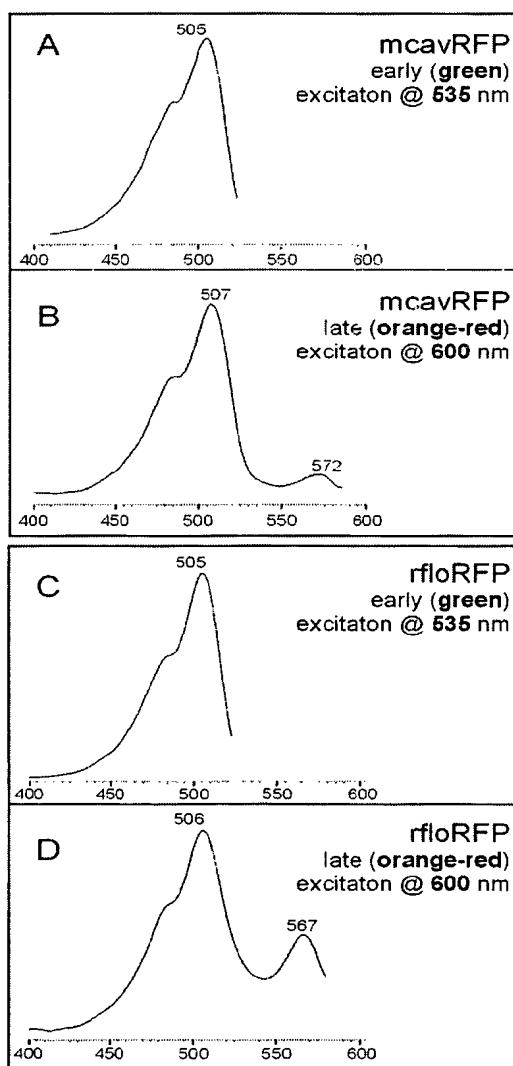
Figure 1.**Figure 2.**

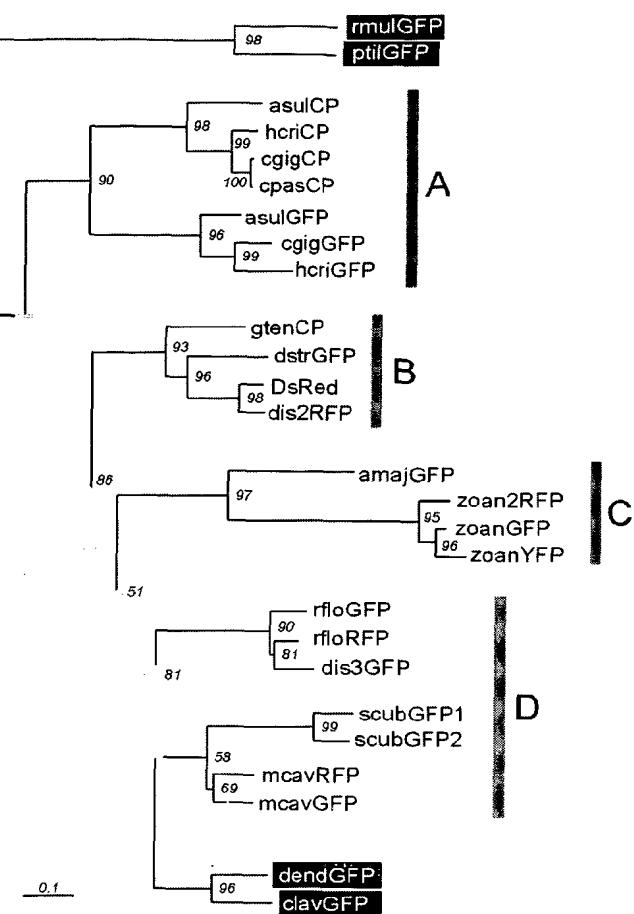
Figure 3.

Figure 4

Protein ID (original ID)	GenBank accession #	Reference	Taxonomy Genus species (Class, Sub-class, Order)	Excitation maxima, nm	Emission maxima, nm	Representative spectra excitation _____ emission _____	Representative spectra excitation _____ emission _____	Color	Representative chromophore structure
amaGFP (amFP486) dstrGFP (dsFP483) clavGFP (CP484)	AF168421 AF168420 AF168424	2 2 2	<i>Anemonia majano</i> (Anthozoa, Zoantharia, Actiniaria) <i>Discosoma striata</i> (Anthozoa, Zoantharia, Corallimorpharia) <i>Clavularia</i> sp. (Anthozoa, Zoantharia, Actiniaria)	458 456 443	486 484 483	458 486 amaGFP	458 486 amaGFP	ORANGE	
GFP cgGFP hcGFP	M62653 AY037776 AF420592	34 this paper this paper	<i>Aequorea victoria</i> (Hydrozoa, ... Hydrozoa) <i>Conchalictus gigantea</i> (Anthozoa, Zoantharia, Actiniaria) <i>Heteroclitus crisia</i> (Anthozoa, Zoantharia, Actiniaria)	395, 471 398, 462 405, 481	508 492 499	395, 471 498 486 cgGFP	395, 471 498 486 cgGFP	GREEN	
ptlGFP rmuGFP zoanGFP (ZFp506) asulGFP (aspF499)	AY015995 AY015996 AF168422 AF532221	35 35 2 4	<i>Philactis sp.</i> (Anthozoa, Alcyonaria, Pennatulacea) <i>Renilla muelleri</i> (Anthozoa, Alcyonaria, Pennatulacea) <i>Zoanthus</i> sp. (Anthozoa, Zoantharia, Zoanthidea) <i>Anemonia sulcata</i> (Anthozoa, Zoantharia, Actiniaria)	500 498 496 493	508 510 508 506	395, 471 498 486 483 zoanGFP	395, 471 498 486 483 zoanGFP	RED	
dis3GFP dendGFP mcavGFP scubGFP1 scubGFP2	AF420593 AF420591 AY037769 AY037772 AY037771	this paper this paper this paper this paper this paper	<i>Discosoma sp. 3</i> (Anthozoa, Zoantharia, Corallimorpharia) <i>Dendrophiliphyia</i> sp. (Anthozoa, Zoantharia, Alcyonacea) <i>Ricordea curvirostra</i> (Anthozoa, Zoantharia, Zoanthidea) <i>Ricordea florida</i> (Anthozoa, Zoantharia, Corallimorpharia) <i>Scolymia eburnea</i> (Anthozoa, Zoantharia, Scleractinia) <i>Scolymia rubens</i> (Anthozoa, Zoantharia, Scleractinia)	503 503 503 503 497	508 510 508 515 506	395, 471 498 486 483 497 zoanYFP	395, 471 498 486 483 497 508 538	YELLOW	
zoanYFP (ZFp538)	AF168423	2	<i>Zoanthus</i> sp. (Anthozoa, Zoantharia, Zanthidea)	494, 528	538	395, 471 498 486 483 508 538 zoanYFP	395, 471 498 486 483 508 538 zoanYFP	?	
DsRed (dFP583) dis2RFP (dsFP593) zoa2RFP	AF168419 AF272711 AY059642	2 36	<i>Discosoma</i> sp. 1 (Anthozoa, Zoantharia, Corallimorpharia) <i>Discosoma</i> sp. 2 (Anthozoa, Zoantharia, Corallimorpharia) <i>Zoanthus</i> sp. 2 (Anthozoa, Zoantharia, Zoanthidea)	558 573 563	583 593 574	395, 471 498 486 483 508 538 zoa2RFP	395, 471 498 486 483 508 538 zoa2RFP	DS RED	
mcavRFP riflRFP	AY037770 AY037773	this paper this paper	<i>Montastraea cavernosa</i> (Anthozoa, Zoantharia, Scleractinia) <i>Ritterellaflorida</i> (Anthozoa, Zoantharia, Corallimorpharia)	507, 512 506, 567	519, 580 516, 574	395, 471 498 486 483 508 538 mcavRFP	395, 471 498 486 483 508 538 mcavRFP	ORANGE-RED	
asulCP (ascCP)	AF246709	3, 4	<i>Anemonia sulcata</i> (Anthozoa, Zoantharia, Actiniaria)	568	none	395, 471 498 486 483 508 538 absorbance only (asulCP)	395, 471 498 486 483 508 538 absorbance only (asulCP)	PURPLE-BLUE	
henICP (hcCP) egICP (egCP) cpasCP (cpCP) genICP (gCP)	AF63776 AF63775 AF383155 AF383156	5 5 5 5	<i>Heriarcis crisia</i> (Anthozoa, Zoantharia, Actiniaria) <i>Conchalictus gigantea</i> (Anthozoa, Zoantharia, Actiniaria) <i>Conchalictus pastiflora</i> (Anthozoa, Zoantharia, Actiniaria) <i>Goniopora temundus</i> (Anthozoa, Zoantharia, Scleractinia)	578 571 571 580	none none none none	395, 471 498 486 483 508 538 absorbance only (asulCP)	395, 471 498 486 483 508 538 absorbance only (asulCP)	?	

Table 1. Summary of spectral features and chromophore structures in the family of GFP-like proteins. Note that this paper uses different names for GFP-like proteins than proposed in original publications (the original names, where available, are given in brackets in the first column; see text for nomenclature details).

Figure 5

Table 2

clade	colors	Zoantharia orders
A	Green, purple-blue	Actiniaria
B	Green, orange-red, purple-blue	Corallimorpharia, Scleractinia
C	Green, yellow, orange-red	Actiniaria, Zoanthidea
D	Green, orange-red	Corallimorpharia, Scleractinia

Figure 6

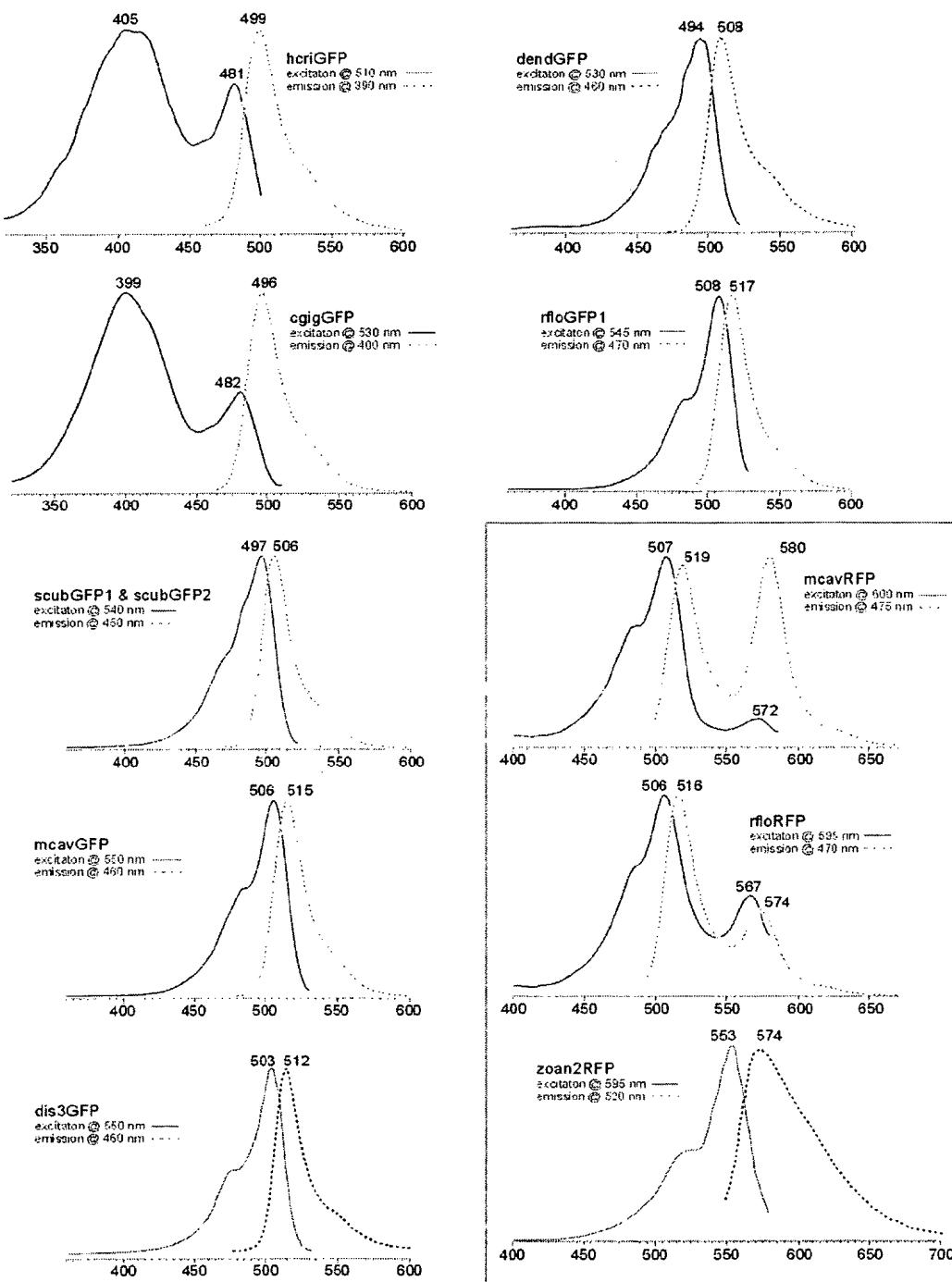


Figure 7

FIGURE 8

Green fluorescent protein from *Heteractis crispa* hcrlGFP

10	20	30	40	50	60
ATTTTGGACAGGTGTTCAACCAAGCAAATTAAAGAAGTCATCATCTTATCTCAGTCAGG					
70	80	90	100	110	120
AAAATGTGTTCTTACATCAAAGAAACCATGCAAAGTAAGGTTACATGGAAGGAAAAGTT					
M	C	S	Y	I	K
E	T	M	Q	S	K
V	Y	M	E	G	K
130	140	150	160	170	180
AACGACCACAACTTCAAGTGCAGTCAGGAAAAGGAGAACCATACAAAGGCTCACAA					
N	D	H	N	F	K
C	T	A	E	G	K
G	E	P	Y	K	G
190	200	210	220	230	240
AGCCTGACGATCACCGTAAGTGAAGGAGGTCTCTGCCATTGCGCTTCGACATTCTTCA					
S	L	T	I	T	V
T	E	G	G	P	L
P	F	A	F	A	F
I	L	S	D	I	L
250	260	270	280	290	300
CACGCCTTCGATATGGCAATAAGGTGTCGCCAAGTACCCCAAAGATCATCCTGATT					
H	A	F	R	Y	G
N	K	V	F	A	K
Y	P	K	D	H	P
E	G	Y	H	P	D
310	320	330	340	350	360
TTTAAGCAGTCTCTCCTGAAGGTTTACTGGGAAAGAGTAAGCAACTATGAGGACGGA					
F	K	Q	S	L	P
K	Q	E	E	G	F
370	380	390	400	410	420
GGAGTCCTTACCGTAAACAAAGAAACTAGTCTGGAGGGAGATTGCATTATTGCAAAATT					
G	V	L	T	V	K
T	Q	E	T	S	L
430	440	450	460	470	480
AAAGCACATGGCACTAACTTCCCCGCAGATGGTCCGGTGATGCAAAACGGACCAATGGA					
K	A	H	G	T	N
F	P	A	D	G	G
490	500	510	520	530	540
TGGGAGCCATCAACTGAAACGTTATTCCACGGGGTGGAGGAATTCTGATGCGCGATGTG					
W	E	P	S	T	E
V	I	P	R	G	G
550	560	570	580	590	600
CCCGCACTGAAGCTGCTTGGTAACAAAGGACATCTCTCTGCGTCATGGAAACAAC					
P	A	L	K	L	L
G	N	K	G	H	L
610	620	630	640	650	660
AAAGTCAAAAAAAGGTGAACCTGCCAAACCGCACTTCATCATTTGAGAATGGAGAAG					
K	S	K	K	G	E
P	A	K	P	H	F
670	680	690	700	710	720
GATAGTGTAGTGACGATGAGAAGACCATTGAGCAGCACGAGAATGTGAGGGCAAGCTAC					
D	S	V	S	D	D
E	K	T	I	E	Q
H	H	H	N	V	R
730	740	750	760	770	780
TTCAATGATAGTGGAAAATGATCATTCTTATTGATTCAATGTTAGGGCATTCAAGTT					
F	N	D	S	G	K
*					
790	800	810	820	830	840
CCAAATTCTTAGACACAGTCTTCCTTAGCTCGTAGCCTACTTACCCATGTTG					
850	860				
TTGAAGTCAATAATAGCTAAGCACTAC (SEQ ID NOS: 01 & 02)					

Figure 9

Green fluorescent protein from *Dendronephthya* sp. dendGFP

10	20	30	40	50	60
5' CATATCGAGAAAGTTGTGAAACCAAATTCTTACTCTACTTTACTACCATGAATCTGATT					
M N L I					
70	80	90	100	110	120
AAAGAAGATATGAGGGTTAAGGTGCATATGGAAGGGAATGTAAACGGGCATGCTTTGTG					
K E D M R V K V H M E G N V N G H A F V					
130	140	150	160	170	180
ATTGAAGGGAAAGGAAAAGGAAGGCCCTACGAAGGGACACAGACCTTGAACCTGACAGTG					
I E G E G K G R P Y E G T Q T L N L T V					
190	200	210	220	230	240
AAAGAAGGCGCGCCTCTCCCATTCTACGACATCTGACAAACAGCATTGCACTACGGA					
K E G A P L P F S Y D I L T T A L H Y G					
250	260	270	280	290	300
AACAGAGTATTCACTGAATACCCAGCAGATATCACGGATTATTCAAGCAATCATTCCCT					
N R V F T E Y P A D I T D Y F K Q S F P					
310	320	330	340	350	360
GAAGGATATTCCCTGGGAAAGAACCATGACTTATGAAGACAAGGGCATTTGTACCATCAGA					
E G Y S W E R T M T Y E D K G I C T I R					
370	380	390	400	410	420
AGCGACATAAGCTTGAAGGTGACTGCTTTCCAAAACATTCTGTTTAATGGGATGAAC					
S D I S L E G D C F F Q N I R F N G M N					
430	440	450	460	470	480
TTTCCCCAAATGGTCCAGTTATGCAGAAGAAAACCTTGAAGTGGGAACCA1CCACAGAG					
F P P N G P V M Q K K T L K W E P S T E					
490	500	510	520	530	540
AAGCTGCACGTGCGTGATGGTTGCTTGTGGTAATATTAACATGGCTCTGCTGCTTGAA					
K L H V R D G L L V G N I N M A L L L E					
550	560	570	580	590	600
GGAGGTGGACATTACCTGTGTGACTTCAAAACACTTACAAAGCGAAGAAGGTTGTCAG					
G G G H Y L C D F K T T Y K A K K V V Q					
610	620	630	640	650	660
TTGCCAGATTATCATTTGTGGACCATCGCATTGAGATCTTGAGTAATGACAGCGATTAC					
L P D Y H F V D H R I E I L S N D S D Y					
670	680	690	700	710	720
AACAAAGTGAAGCTGTACGAGCATGGGGTTGCTCGTATTCTCCGTTGCCAAGTCAGGC					
N K V K L Y E H G V A R Y S P L P K S G					
730	740	750	760	770	780
CTGGTAGAGGTTCAAGGGAAAGCCATAATGACTGCATAGATAAACATGTAGTGAAGACCA					
L V E V Q G K A I M T A *					
790	800	810	820	830	840
CATACTCGGGATTAGAGTTAGGGATTGGTAGTTGTGGTAGATTCTAGCCTACAAATT					
TTGGG 3' (SEQ ID NO:03 & 04)					

Figure 10
Red fluorescent protein from *Zoanthus* sp. zoanRFP

10	20	30	40	50	60
GAGTTGAGTTCTCGACTTCAGTTGTATCACITTGACGTATCAAGTGATCTATTCTCAAC					
70	80	90	100	110	120
ATGGCCCATTCAAAGCACGGACTAACAGATGACATGACAATGCATTCCGTATGGAAGGG					
M	A	H	S	K	H
G	L	T	D	D	M
D	M	T	M	H	F
R	M	E	G		
130	140	150	160	170	180
TGCCTCGATGGACATAAGTTGTAATCGAGGGCAACGGCAATGAAATCCTTCAAAGGG					
C	V	D	G	H	K
F	V	I	E	G	N
G	N	G	N	G	N
P	F	K	F	K	G
190	200	210	220	230	240
AAACAGTTTATTAATCTGTGTGATTGAAGGGAGGACACTGCCATTCTCCGAAGACATA					
K	Q	F	I	N	L
C	V	I	E	G	G
P	L	P	F	S	E
L	P	F	S	E	D
250	260	270	280	290	300
TTGTCTGCTCGTTGACTACGGAAACAGGCTCTTCACTGAATATCCTGAAGGCATAGTT					
L	S	A	A	F	D
Y	G	N	R	L	F
G	N	R	L	F	T
E	P	E	T	E	Y
G	E	R	P	P	P
I	V	S	F	E	G
V	S	C	A	G	I
310	320	330	340	350	360
GACTATTCAAGAACACTCGTGTCTGCTGGATATACGTGGCACAGGTCTTCGCTTGAA					
D	Y	F	K	N	S
Y	F	K	N	S	C
E	K	N	S	C	P
A	N	S	C	P	A
G	R	R	S	C	G
370	380	390	400	410	420
GATGGAGCAGTTGCATATGCAGTCAGATATAACAGTAAATGTTAGGGAAAAGTCATT					
D	G	A	V	C	I
G	A	V	C	I	C
A	C	I	S	A	S
T	T	I	C	S	A
D	I	T	T	V	N
I	T	V	N	V	R
430	440	450	460	470	480
TATCATGAGTCCACGTTTATGGAGTGAACCTTCCTGCTGATGGACCTGTGATGAAAG					
Y	H	E	S	T	F
H	E	S	T	F	Y
E	S	T	F	Y	G
S	T	F	Y	G	V
T	F	Y	G	V	N
F	P	A	P	A	F
490	500	510	520	530	540
ATGACAACATAATTGGGAACCGCTCCTGCGAGAAAATCATACCAATAATAGTCAGAAAGATA					
M	T	T	N	W	E
T	T	N	W	E	P
N	W	E	P	S	C
W	E	P	S	C	E
E	P	S	C	E	K
550	560	570	580	590	600
TTAAAAGGGGATGTCCTCATGTACCTCCTCTGAAGGATGGTGGCGTTACCGCTGCCAG					
L	K	G	D	V	S
K	G	D	V	S	M
G	D	V	S	M	Y
D	V	S	M	Y	L
V	S	M	Y	L	L
S	M	Y	L	L	K
M	Y	L	L	K	D
610	620	630	640	650	660
TTTGACACAAATTACAAAGCAAAGACTGAGCCAAAAGAAAATGCCGGACTGGCACTTCATC					
F	D	T	I	Y	K
D	T	I	Y	K	A
T	I	Y	K	A	K
I	Y	K	A	K	T
670	680	690	700	710	720
CAGCATAAGCTCAACCGTGAAGACCGCAGCGATGCTAAGAATCAGAAATGGCAACTGATA					
Q	H	K	L	N	R
H	K	L	N	R	E
K	L	N	R	E	D
L	N	R	E	D	R
N	R	E	D	R	S
R	E	D	R	S	D
E	D	R	S	D	A
D	R	S	D	A	K
R	S	D	A	K	N
S	A	D	K	N	Q
A	L	A	K	Q	K
L	P	L	M	P	D
P	*	P	P	D	W
*	P	P	E	D	H
730	740	750	760	770	780
GAACATGCTATTGCATCCGATCTGCTTACCTGATAACAAAGGAGTTGCTATTGCATG					
E	H	A	I	A	S
H	A	I	A	S	R
A	I	A	S	R	S
I	A	S	R	S	A
S	R	S	A	L	P
R	S	A	L	P	*
790	800	810	820	830	840
TGCATGCCTATTACGCTGATAAAAAATGTTTTAACATGCAATTGTATGTGCATGCACA					
850					
(SEQ ID NOS:05 & 06)					

Figure 11

Green fluorescent protein from *Scolymia cubensis* scubGFP1 (AY037767)

10 20 30 40 50 60
 5 'TGTGACATTCAAGTCATATAGGAGCCTCTATCGGAGCTGAGGTCCCATTACCGTTGTGAT
 70 80 90 100 110 120
 TTGGACGGGAGCAGATCGAGAACACMAGGGCTGTACGAGTCGATAATTACTTTACAT
 130 140 150 160 170 180
 CTACCAACATGCAGCGTCTGGGATGAAGGTTAAGGAACATATGAAGATCAAACGCGTA
 M Q R A G M K V K E H M K I K L R M
 190 200 210 220 230 240
 TGGGAGGTACTGTAACCGAAAGCATTGCGGTTAATGGGACAGGAGACGGCTACCCCT
 G G T V N G K H F A V N G T G D G Y P Y
 250 260 270 280 290 300
 ATCAGGGAAAACAGATTGAAACTTATCGTCAAGGCAGCGAACCTCTGCCTTCGCTT
 Q G K Q I L K L I V E G S E P L P F A F
 310 320 330 340 350 360
 TTGATATCTTGTCAAGCAGCATTCCAGTATGGCAACAGGCATTACCGAATACCAACAG
 D I L S A A F Q Y G N R A F T E Y P T E
 370 380 390 400 410 420
 AGATAGCAGACTATTCAGCAGTCGTTGAGTTGGCGAGGGGTTCTCCTGGAACGAA
 I A D Y F K Q S F E F G E G F S W E R S
 430 440 450 460 470 480
 GTTTCACCTTCGAAGATGGGCCATTGCGTCGCCACCAACGATATAACGATGGTTGGTG
 F T F E D G A I C V A T N D I T M V G G
 490 500 510 520 530 540
 GTGAGAGTCAGTATGATATTGATTGATGGCTGAACTTCCCTGAAGATGGTCCAGTGA
 E F Q Y D I R F D G L N F P E D G P V M
 550 560 570 580 590 600
 TGCAAAAGAAAACGTAAGGACATCCACTGAGATAATGTATATGCAAAATGGAG
 Q K K T V K W E P S T E I M Y M Q N G V
 610 620 630 640 650 660
 TGCTGAAGGGTGAGGTAAACATGGCTCTGTGCTTCAGACAAAAGCCATTACCGTTGCG
 L K G E V N M A L L Q D K S H Y R C D
 670 680 690 700 710 720
 ACCTCAAAACTACTAACAAAGCTAAAGATAATGTGCCCATCCTCCAGGCTACCACTATG
 L K T T Y K A K N N V P H P P G Y H Y V
 730 740 750 760 770 780
 TGGATCACTGCATTGAAATACTCGAAGAACGTAAGGATCACGTTAACGCTGCGGAGCATG
 D H C I E I L E R K D H V K L R E H A
 790 800 810 820 830 840
 CTAAAGCTCGTTCTAGCCTGTCACCTACCAGTGCAAAAGAACGAAAGGTTAGGTGATAG
 K A R S S L S P T S A K E R K A *
 850 860 870 880 890 900
 TCAAAAGACAACAAAGACGAAAATGAAAGGTGTCATTGTTAGAATTGATATTCGAT
 910 920 930 940 950 960
 TCAATGATTGCTTAAGGGATTGCTAGAGGCTAGCTAACAGGTTAACATCATAGGATAG
 970 980 990 1000 1010 1020
 AGATTYCCTGCGGAGTTAGAACCTTATTTCCGAAATTCCACMCTAGAGTCGTTGAGA
 1030 1040 1050 1060 1070 1080
 AATTTATAGAGACTAGCTTAGAGTTACTTTGCGAAAAAAAGGTTCCATTGTC
 1090 1100 1110 1120 1130 1140
 GTTATTACAGCATTAAAGCATAGGAATAGAGATTGCGTTATGGAAAATAACAGTAGGAA
 1150 1160 1170
 AATACTGTTGAAAATAACTTGTGTCGAAAAAAA 3 '

(SEQ ID NOS:07&08)

FIGURE 12

Green fluorescent protein from *Scolymia cubensis* scubGFP2 (AY037771)

10 20 30 40 50 60
 5 ' CCTGGTGATTGGACGAGAGCAGATCGAGAAATAGCAAGGTTTACCAAGCGTGATAATTAA
 70 80 90 100 110 120
 CTTTACATCTAACACATGCAATCTGCTGGAGAAGAATGTCGTTAAGGACTTCATGAA
 M Q S A G K K N V V K D F M K

 130 140 150 160 170 180
 GATCACACTGCGTATGGACGGTGCCTAAACGGGAAGGCCCTCGCGGTTAATGGAACAGG
 I T L R M D G A V N G K P F A V N G T G

 190 200 210 220 230 240
 AGATGGCAACCCCTTATGGTGGAAATACAGACTTGAAGCTTACCGTCGATGGCAACAAACC
 D G N P Y G G I Q S L K L T V D G N K P

 250 260 270 280 290 300
 TCTGCCTTTGCTTTGATATCTGTCAGCAGCATTCCAGTATGGCAACAGGGCATTCAC
 L P F A F D I L S A A F Q Y G N R A F T

 310 320 330 340 350 360
 CGAATACCCAAAAGAGATATCAGACTATTCAAGCAGTCGTTGAGTTGGCGAGGGTT
 E Y P K E I S D Y F K Q S F E F G E G F

 370 380 390 400 410 420
 TACCTGGGAACGAAGTTCACTTCAAGACGGGGCATTGCGTCGCCACGAACGATAT
 T W E R S F T F E D G A I C V A T N D I

 430 440 450 460 470 480
 AAAGATGGTGGCGATGAGTTCAATATAACATTGATTGATGGTGTGAATTCCCTGA
 K M V G D E F Q Y N I R F D G V N F P E

 490 500 510 520 530 540
 AGATGGTCWGTYATGCAGAAGAAAACGGTGAAGTGGGAGGCATCCACAGAGATAATGCG
 D G P V M Q K K T V K W E P S T E I M R

 550 560 570 580 590 600
 TGTGCAAGGTGGAGTGCTAAAGGGTGAAGGTTAACATGGCTGTGCTTAAAGACAAAAG
 V Q G G V L K G E V N M A L L L K D K S

 610 620 630 640 650 660
 CCATTACCGATGTGACTTCAAAACTACTTACAAAGCTAAGAAATCCTGTCCCGCCGACGGC
 H Y R C D F K T T Y K A K N P V P P T A

 670 680 690 700 710 720
 GCTTCAGACTACCACTATGGATCACTGTATTGAAATCACCGAGGAAAATAGGGATTA
 L P D Y H Y V D H C I E I T E E N R D Y

 730 740 750 760 770 780
 CGTTAAGCTGCAGGAGTATGCTAAAGCTCGTCTGGCCTGCACCTGCCCGAACTGCAAAA
 V K L Q E Y A K A R S G L H L P E L Q K

 790 800 810
 GTAAAGGCTTAGGCGATAGTCAGACGACAACGAGAAGA 3'
 *

(SEQ ID NO:09 & 10)

FIGURE 13

Red fluorescent protein from *Ricordia florida* rflcRFP (AY037773)

10 20 30 40 50 60
 5' TGTGAAAGTTAACATTTACTTACTTCTACCAGCATGAGTGCACCAAAGAGGAAATGA
 M S A L K E E M K

 70 80 90 100 110 120
 AAATCAAGCTTACATGGTGGCGTTGTTAACGGGCACCCATTCAAGATCATTGGGACG
 I K L T L V G V V N G H P F K I I G D G

 130 140 150 160 170 180
 GAAAAGGCAACCCCTATGAGGGATCGCAGGAATTAACCCCTGCCGTGGTGAAGGAGGC
 K G K P Y E G S Q E L T L A V V E G G P

 190 200 210 220 230 240
 CTCTGCCTTCTCTTATGATATCCTGACAACGATAGTTCACTATGGCAACAGGGCATTG
 L P F S Y D I L T T I V H Y G N R A F V

 250 260 270 280 290 300
 TGAACTACCCAAAGGACATACCAGATATTTCAAGCAGACCTGCTCTGGTCCCTGGTCTG
 N Y P K D I P D I F K Q T C S G P G A G

 310 320 330 340 350 360
 GATATTCCTGGCAAGGACCATGAGTTGAAAGACGGAGGCCCTGCAGCTACGAGCC
 Y S W Q R T M S F E D G G V C T A T S H

 370 380 390 400 410 420
 ATATCAGGGTGGATGGCAGACTTCAATTATGACATTCACTTCATGGGAGCGGATTTC
 I R V D G D T F N Y D I H F M G A D F P

 430 440 450 460 470 480
 CTCTTAATGGTCCAGTGATGCGAGAAAAGAACAGTGAAATGGGAGCCATCCACTGAGATAA
 L N G P V M Q K R T V K W E P S T E I M

 490 500 510 520 530 540
 TGTTTCAATGTGATGGATTGCTGAGGGGTGATGTTGCCATGTCCTGTTGCTGAAAGGAG
 F Q C D G L L R G D V A M S L L K G G

 550 560 570 580 590 600
 GCGGCCATTACCGATGTGACTTTAAACTATTTATAAACCCAAAGAAGAATGTCAAGATGC
 G H Y R C D F K T I Y K P K K N V K M P

 610 620 630 640 650 660
 CAGGTTACCATTTGAGGACACTGCATTGAGATAACGAGTCAACAGGACGATTACAACG
 G Y H F V D H C I E I T S Q Q D D Y N V

 670 680 690 700 710 720
 TGTTGAGCTGTACGAGGGTGCTGAGCCCACACTCTCCTCTGCAGAAACCATGCAAG
 V E L Y E G A V A H Y S P L Q K P C Q A

 730 740 750 760 770 780
 CAAAGGCATAAACGCAAACAAACCAAGAGGACAACAAGACATTAAATCAAATCACATCTT
 K A *

790 800
TGTATTTGGTTAGAGTTGAAAAAAA 3'

(SEQ ID NO:11 & 12)

FIGURE 14

Green fluorescent protein from *Ricordea florida* rfloGFP (AY037772)

10 20 30 40 50 60
 5 'AGTCACCTCGGTGTTTTAGGACAGGAAGGATCACGAGCAAGAGAAGAACTGTGAAAGTT
 70 80 90 100 110 120
 AACACTTTACTCTACTTCTACCACGATGAGTGCACCAAAGAGGAAATGAAATCAAGCT
 M S A L K E E M K I K L

 130 140 150 160 170 180
 TAAAATGGTGGCGTTGTTAACGGCAGTCATTTCAGATCGATGGGAAGGAAAAGGCAA
 K M V G V V N G Q S F Q I D G E G K G K

 190 200 210 220 230 240
 ACCTTACGAGGGATCACAGAAATTAACCCCTGAAAGTGGTGGAAAGGAGGGCCTCTGCTCTT
 P Y E G S Q K L T L E V V E G G P L L F

 250 260 270 280 290 300
 CTCTTATGATATCCTGACAACGATATTCAGTATGCCAACAGGGCATTCTGAACTACCC
 S Y D I L T T I F Q Y G N R A F V N Y P

 310 320 330 340 350 360
 AAAGGACATACCAAGATATTTCAAGCAGACCTGCTCTGGTCTGATGGTGGATTTCCTG
 K D I P D I F K Q T C S G P D G G F S W

 370 380 390 400 410 420
 GCAAAGGACCATGACTTATGAAGACGGAGGGTTGCACTGCTCAAACACATCAGCGT
 Q R T M T Y E D G G V C T A S N H I S V

 430 440 450 460 470 480
 GGACGGCGACACTTTTATTATGTGATAAGATTAATGGAGAGAATTTCCCTCAAATGG
 D G D T F Y Y V I R F N G E N F P P N G

 490 500 510 520 530 540
 TCCAGTAATGCAGAAAAGAACAGTGAAATGGGAGCCATCCACTGAGATAATGTTCAACG
 P V M Q K R T V K W E P S T E I M F E R

 550 560 570 580 590 600
 TGATGGATTGCTGAGGGGTGACATTGCCATGCTCTGTTGCTGAAAGGAGGCAGGCCATTA
 D G L L R G D I A M S L L K G G G H Y

 610 620 630 640 650 660
 CCGATGTGACTTTAAACTATTATACACCCAAAGAGGAAGGTCAACATGCCAGGTTACCA
 R C D F K T I Y T P K R K V N M P G Y H

 670 680 690 700 710 720
 TTTTGTGGACCACTGCATTGAGATAACAGAAGCAGCACAAGGATTACAAACATGGCTGTGCT
 F V D H C I E I Q K H D K D Y N M A V L

 730 740 750 760 770 780
 CTCTGAGGATGCTGTAGCCCCACAACCTCTCTGAGAAAAAAAGCCAAGCAAAGGCGTA
 S E D A V A H N S P L E K K S Q A K A *

 790
 AAGCCAAACAAACCTAA 3'

(SEQ ID NO:13&14)

Figure 15

Red fluorescent protein from *Montastraea cavernosa* mcavRFP (AY037770)

```

  10          20          30          40          50          60
5' ACGCAGGGATTCACCTGGTGATTGGAAGAGAGCAGACCGAGAACAAAGAGCTGTAT
  70          80          90          100         110         120
AAGGCTGATATCTTACGTCTACCATCATGAGTGTGATTAATCAGTCATGAAGAT
R L I S Y F T S T I M S V I K S V M K I

  130         140         150         160         170         180
CAAGCTCGCTATGGAAGGCAGTGTAAACGGGCACAACCTCGTAATTGTTGGAGAAGGAGA
K L R M E G S V N G H N F V I V G E G E

  190         200         210         220         230         240
AGGCAAGCCTTATGAGGGAACACAGAGTATGGACCTTACAGTCAGAACAGAGGCACCTCT
G K P Y E G T Q S M D L T V K E G A P L

  250         260         270         280         290         300
GCCTTTCGCTACGATATCATGACAACAGTATTCCATTACGGCAATAGGGTATTGCAAA
P F A Y D I M T T V F H Y G N R V F A K

  310         320         330         340         350         360
ATACCCAAAACATATCCCAGACTATTCAAGCAGATGTTCCCTGAGGACTATTCCGGAA
Y P K H I P D Y F K Q M F P E E Y S W E

  370         380         390         400         410         420
ACGAAGCATGAATTTCGAAGGGGGGCATTGACCCAGGAACAGAGATAACAATGGAA
R S M N F E G G I C T A R N E I T M E

  430         440         450         460         470         480
AGGCAGACTGTTTTTCATAAAAGTTCGATTGATGGTGTGAACTTCCCCCAATGGTCC
G D C F F N K V R F D G V N F P P N G P

  490         500         510         520         530         540
AGTCATGCAAGAAGAACCGCTGAAATGGAGGCCATCCACTGAAAAATGTATGCGTGA
V M Q K K T L K W E F S T E K M Y V R D

  550         560         570         580         590         600
TGGAGTGTGACGGGTGATATCAACATGGCTTTGTTGAGGAGGTGGCATTACCG
G V L T G D I N M A L L E G G G H Y R

  610         620         630         640         650         660
ATGTGACTTCAGAACTACTTACAGAGCTAAGAAGAAGGGTGTCAAGTTACCAAGATTATCA
C D F R T T Y R A K K K G V K L P D Y H

  670         680         690         700         710         720
CTTTGAGGATCACTCCATTGAGATTTGCGCCATGACAAAGAACACTGAGGTTAACGCT
F E D H S I E I L R H D K E Y T E V K L

  730         740         750         760         770         780
GTATGAGCATGCCGAAGCTCATTCTGGCTGCCGAGGGTGGCAAAGTAAAGGCTAACGA
Y E H A E A H S G L P R V A K *

  790
AAAGCCAAGACCACA 3'

```

(SEQ ID NO:15 & 16)

FIGURE 16

Green fluorescent protein from *Montastraea cavernosa* mcavGFP (AY037769)

10 20 30 40 50 60
 5' ATTCGCCCCTGGTGAATTGGAAGAGAGCAGATCGAGAACAAACAGAGCTGTAAGGTTGATA
 70 80 90 100 110 120
 TCTTACTTACGTCTACCATCATGACAAGTGTGACAGGAAAAGGGTGTGATTAACCCAG
 M T S V A Q E K G V I K P D

 130 140 150 160 170 180
 ACATGAAGATGAAGCTCGTATGGAAGGTGCTGTAACGGGCACAAGTTCGTGGTTGAAG
 M K M K L R M E G A V N G H K F V V E G

 190 200 210 220 230 240
 GAGATGGAAAAGGGAAGGCCTTCGACGGAACACAGACTATGGACCTTACAGTCATAGAAG
 D G K G K P F D G T Q T M D L T V I E G

 250 260 270 280 290 300
 GCGCACCATTCGCTTACGATATCTGACAAACAGTATTGATTACGGCAACAGGG
 A P L P F A Y D I L T T V F D Y G N R V

 310 320 330 340 350 360
 TATTGCGCAAATACCCAGAACAGACATAGCAGATTATTCAGAACAGCTTCTGAGGGGT
 F A K Y P E D I A D Y F K Q T F P E G Y

 370 380 390 400 410 420
 ACTTCTGGAACGAAAGCATGACATACGAAGAACAGGGCATTGATCGCCACAAACGACA
 F W E R S M T Y E D Q G I C I A T N D I

 430 440 450 460 470 480
 TAACAATGATGGAAGGCCTCGACGACTGTTGCTATAAAATTGATTTGATGGTGTGA
 T M M E G V D D C F A Y K I R F D G V N

 490 500 510 520 530 540
 ACTTTCTGCCAATGGTCAGTTATGCAGAGGAAGACGCTGAATGGGAGCCATCCACTG
 F P A N G P V M Q R K T L K W E P S T E

 550 560 570 580 590 600
 AGATAATGATGCGCCTGATGGAGTGCTGAAGGGTGTAAACATGGCTCTGTTGCTTG
 I M Y A R D G V L K G D V N M A L L L E

 610 620 630 640 650 660
 AAGGAGGTGGCCATTACCGATGTGACTTCAAAACTACTTACAAAGCTAAGAAGGTTGTCC
 G G H Y R C D F K T T Y K A K K V V R

 670 680 690 700 710 720
 GGTTGCCAGACTATCACTTGTGGACCATCGCATTGAGATTGTGAGCCACGACAAAGATT
 L P D Y H F V D H R I E I V S H D K D Y

 730 740 750 760 770 780
 ACAACAAGGTTAACGTCACGAGCATGCCAAGCTCGTCATGGACTGTCAAGGAAGGCCA
 N K V K L H E H A E A R H G L S R K A K

 790 800 810 820 830 840
 AGTAAAGGCTTAATGAAAGTCAGACGACAACGAGGAGAACAAAGTACTTTTGTTA
 *

 850 860 870 880 890 900
 AATTGAAAGGCATTTACTCGGAATTAGTAGATTGATACTTTGATTCAAGGATTGTTCCG
 910 920 930 940 950 960
 GGATTGTTAGAGACTAGCTCTAGAGTTGTATTTGTGAAAAAAAGATAGTTCCAGTTT
 970 980 990 1000 1010 1020
 TGCAGGGATTACAGCATGGGGATAGACTTTAAACTCAGTTGTGGTCAAATGCAAGTAAG
 1030 1040 1050 1060
 AAAACTGTAGTGAGAATAAAACTTGTATCGAAGCCGAAAAAA 3'

(SEQ ID NOS: 17 & 18)

Figure 17

Green fluorescent protein from *Condylactis gigantea* cgigGFP (AY037776)

10	20	30	40	50	60
5' ACAGCTGTTCATCCACGCTCATTCAAGACGCCGTCAACTTATTCCAGTCAGGAAAATGT					
M Y					
70	80	90	100	110	120
ATCCTTGGATCAAGGAAACCATGCGCAGTAAGGTTTACATGGAAGGAGATGTTAACACC					
P W I K E T M R S K V Y M E G D V N N H					
130	140	150	160	170	180
ACGCCCTCAAGTGCACTGCAGTAGGAGAAGGAAACCATACAAAGGCTCACAGACCTGA					
A F K C T A V G E G K P Y K G S Q D L T					
190	200	210	220	230	240
CGATTACCGTCACTGAAGGAGGTCTCTGCCATTGCTTTGACATTCTTCACACGCC					
I T V T E G G P L P F A F D I L S H A F					
250	260	270	280	290	300
TTCAGTATGGCAACAAAGGTGTTACCGATTACCCGACGATATTCTGATTTCTTAAGC					
Q Y G N K V F T D Y P D D I P D F F K Q					
310	320	330	340	350	360
AGTCTCTCTGGATGGTTTACTGGAGAAGAGTAAGCACSTATGACGATGGAGGAGTC					
S L S D G F T W R R V S T Y D D G G V L					
370	380	390	400	410	420
TCACAGTTACCCAAGACACTAGTCTGAAGGGAGATTGCATTATTGCAACATTAAAGTC					
T V T Q D T S L K G D C I I C N I K V H					
430	440	450	460	470	480
ATGGCACTAACTTCCCGAAATGGTCCGGTGTGCAAACAAAGACCGATGGATGGAGC					
G T N F P E N G P V M Q N K T D G W E P					
490	500	510	520	530	540
CATCCACCACTGAAACGGTTATTCCACAAGATGGAGGAATTGTTGCTGCGCGATACCCG					
S S T E T V I P Q D G G I V A A R S P A					
550	560	570	580	590	600
CACTAAGGCTGCGTGATAAAGGTCATTTATCTGCCACATGGAAACAACTTACAAGCAA					
L R L R D K G H L I C H M E T T Y K P N					
610	620	630	640	650	660
ACAAAGAGGTGAAGCTGCCAGAACCTCCACTTTCATCATTTGCGAATGGAAAGCTGAGTG					
K E V K L P E L H F H H L R M E K L S V					
670	680	690	700	710	720
TTAGTGACGATGGGAAGACCATTAAAGCAGCACCGAGTATGTGGTGGCTAGCTACTCAAAG					
S D D G K T I K Q H E Y V V A S Y S K V					
730	740	750	760	770	780
TGCCTTCGAAGATAGGACGTCAATGATCATTTCCCTTATTAAATATCAATGATGTGGCTT					
P S K I G R Q *					
790	800	810	820	830	840
TCAATTTCAAAATTGGTTAACATAGGTCTTTGGATTTGGTAACCCCAACCTT					
850	860	870	880	890	
AATTCCCAATAATTGGAAAGTCAAATAAACAGCCTCCCTGGCCTTAA 3'					

(SEQ ID NOS: 19 & 20)

FIGURE 18

Green fluorescent protein from *Agaricia fragilis* afraGFP (AY037765)

10 20 30 40 50 60
 5' CAAGGAAGCAAATCTTTACCAAGAGATCTCGCGTCAAAGCAACCTATGAGTGATGGCGA
 M A I
 70 80 90 100 110 120
 TTTCTACTCTAAAGAACGTCATCATCATCGTTATTATATACTCCTGCAGCAGTGTGCTG
 S T L K N V I I V I I Y S C S T C A V
 130 140 150 160 170 180
 TTTGGCTCGAATTCAAACCTCTGAATCCTCTTCACTAATGGGATTGCAGAGGAATGAAGA
 W S N S N S E S S F T N G I A E E M K T
 190 200 210 220 230 240
 CTAGGGTACATTTGGAGGGTACTGTTAACGGGCACTCCTTACAATTAAAGGCGAAGGAA
 R V H L E G T V N G H S F T I K G E G R
 250 260 270 280 290 300
 GAGGCTACCCCTACAAAGGAGAACAGTTATGAGCCTTGAGGTCGTCAATGGTGCCTC
 G Y P Y K G E Q F M S L E V V N G A P L
 310 320 330 340 350 360
 TGCCGTTCTCTTTGATATCTTGACACCAGCATTATGTATGGCAACAGAGTGTTCACCA
 P F S F D I L T P A F M Y G N R V F T K
 370 380 390 400 410 420
 AGTACCCACCAAAACATACCAAGACTATTTCAAGCAGACGTTTCTGAAGGGTATCACTGGG
 Y P P N I P D Y F K Q T F P E G Y H W E
 430 440 450 460 470 480
 AAAGAAACATTCCCTTGAAAGATCAGGCCGTGCACGGTAACCGCACATAAGATGG
 R N I P F E D Q A A C T V T S H I R L E
 490 500 510 520 530 540
 AAGAGGAAGAGAGGCCCTTGAAATAACGTCAGATTCACTGTGTGAACCTTCCCCCTA
 E E E R R F V N N V R F H C V N F P P N
 550 560 570 580 590 600
 ATGGTCCAGTCATGCAGAGGAGGATACTGAAATGGGAGGCCACTGAGAACATTATC
 G P V M Q R R I L K W E P S T E N I Y P
 610 620 630 640 650 660
 CGCGTGATGGTTCTGGAGGCCATGTTGATATGACTCTCGGGTTGAAGGAGGTGGCT
 R D G F L E G H V D M T L R V E G G G Y
 670 680 690 700 710 720
 ATTACCGAGCTGAGTCAAAAGTACTTACAAAGGGAAGACCCAGTCCGGACATGCCAG
 Y R A E F K S T Y K G K T P V R D M P D
 730 740 750 760 770 780
 ACTTCACTTCATAGACCAACCGCATTGAGATTACGGAGCATGACGAAGACTACACCAATG
 F H F I D H R I E I T E H D E D Y T N V
 790 800 810 820 830 840
 TTGAGCTGCATGACGTATCTGGCTCGTTACTCTATGCTGCCACTATGTAAGCGGAAA
 E L H D V S W A R Y S M L P T M
 850 860 870 880 890 900
 AGGCAAGGCAACAAGACGCAAAACGCCCTGTTGTCTCTTTCATAAGAGATTGACAA
 910 920 930 940 950 960
 CCGTGGTTCTTGCCATTAAATTGAATTAGTTAAATTAAATCTTGGGATTGATGTAG
 970 980 990 1000 1010 1020
 ACGCTTGGTTGCTAAGTAAGAAAACATTGTGATTATTAATTGTTGCCTGAAGCAA
 1030
 AAAAAAAA 3'

(SEQ ID NOS:21 & 22)

FIGURE 19

Green fluorescent protein from *Ricordia florida* rfloGFP2 (AY037774)

10	20	30	40	50	60
5' AGCCACTTCGGTGTCTGTCGAGAGGAAGGATCACGAACAAGAGAAGAGCTGTAAAAGTT					
70	80	90	100	110	120
AAAATTTACTTACTTCCAGCATGAATGCACTTCAAGAGGAAATGAAAATCAAGCT					
M	N	A	L	Q	E
130	140	150	160	170	180
TACAATGGTGGCGTTGTTAACGGCAGTCATTAAGATCGATGGGAAAGGAAAAGGGAA					
T	M	V	G	V	V
190	200	210	220	230	240
ACCTTACGAGGGATCACAGGAATTGACCCCTAAAGTGGTGGAAAGGCAGGGCTCTGCTCT					
P	Y	E	G	S	Q
250	260	270	280	290	300
CTCTTATGATATCCTGACAAACGATATTCAGTATGGCAACAGGGCATTCTGAACTACCC					
S	Y	D	I	L	T
310	320	330	340	350	360
AAAGGACATACCAAGATATTTCAAGC AACGTGTTCTGGTCTTGATGGCGGATATTCTG					
K	D	I	P	D	I
370	380	390	400	410	420
GCAAAGGACCATGACTTATGAGGACGGAGGGTTTGACTGCTACAAGCAACGTCAGCGT					
Q	R	T	M	T	Y
430	440	450	460	470	480
GGTCGGCAGACACTTCATTATGAAATTCACTTATGGGGCGAATTTCCTCAAATGG					
V	G	D	T	F	N
490	500	510	520	530	540
TCCRG TGATGCAGAAAAGAACAGTGAAGTGGGAGCCCTCCACTGAGATAATGTTGAACG					
P	V	M	Q	K	R
550	560	570	580	590	600
TGATGGATTGCTGAGGGGTGATGTTCCCATGTCCTGTTGCTGAAAGGAGGCGACCATTA					
D	G	L	L	R	G
610	620	630	640	650	660
CCGATGTGACTTTAAACATTTATAAACCCAACAAGAAGGTCAAGCTGCCAGGTTACCA					
R	C	D	F	K	T
670	680	690	700	710	720
TTTGAGGACACTGCATTGAGATAAAAGAGTCAGAGAATGATTACAACATGGTTGCGCT					
F	V	D	H	C	I
730	740	750	760	770	780
CTTTGAGGATGCTGAGCACACTACTCTCTGGAGAAAAGAGCCAGGCCAAAGGCGTA					
F	E	D	A	V	A
790	800	810	820	830	840
AATCCAAACACCTAACAGAACGACAAGGCATTCAATCTAACGATGTTGAATTGG					
850	860	870	880	890	900
GTTAGGAATGTGTTGGGTCAAGACTAGGTCTAGAACGTTTCATTTGGCTGGATTTGTTT					
910	920	930	940	950	960
ACTCAGTTATAGACAAGAAAAAAATCTTAAATGACTTGGGTTGGATTAGCTTCCGGCAC					
970	980	990	1000	1010	1020
TGTCAATTCCGGATTCCCTAGAAATTTGAGACCAAGCCTTTTGAGCTGAGAACGT					
AATC	3'				

(SEQ ID NOS: 23 & 24)

FIGURE 20

Green fluorescent protein from *Montastraea cavernosa* mcavGFP2 (AY037768)

10 20 30 40 50 60
 5 ' AGAGCTGTAGGGTATCTTACGTCTACCATCATGACCAGTGTGCACAGGAAAA
 M T S V A Q E K

 70 80 90 100 110 120
 GGGTGTGATTAACAGACATGAAGATGAAGCTGCATGGAAAGGTGCTGAAACGGGCA
 G V I K P D M K M K L R M E G A V N G H

 130 140 150 160 170 180
 CAAGTTCTGTGATTGAAGGGAGATGGAAAAGGGAAGCCTTCGACCGAACACAGACTATGGA
 K F V I E G D G K G K P F D G T Q T M D

 190 200 210 220 230 240
 CCTTACAGTCATAGAAGGCGCACCATTCGCTTCGCTTACGCTATCTGACAACAGTATT
 L T V I E G A P L P F A Y A I L T T V F

 250 260 270 280 290 300
 CGATTACGGCAACAGGGTATTCGCAAATACCCAGAACAGACATAGCAGATTATTCAAGCA
 D Y G N R V F A K Y P E D I A D Y F K Q

 310 320 330 340 350 360
 GACATTTCCGTGAGGGGACTTCTGGAACGAAAGCATGACATACGAAGAACAGGGCATTG
 T F P E G Y F W E R S M T Y E D Q G I C

 370 380 390 400 410 420
 CATGCCACAAACGACATAACAATGATGAAAGGGCGTCGACGACTGTTTGTCTATAAAAT
 I A T N D I T M M K G V D D C F V Y K I

 430 440 450 460 470 480
 TCGATTGATGGTGTAACTTCTGCCATGGTCCAGTTATGCCAGAGGAAGACGCTGAA
 R F D G V N F P A N G P V M Q R K T L K

 490 500 510 520 530 540
 ATGGGAGCCATCCACTGAGAAAATGTATGCCGTGATGGAGTGTGAAGGGTGTAA
 W E P S T E K M Y A R D G V L K G D V N

 550 560 570 580 590 600
 CATGGCTCTGTTGCTGAAGGAGGTGGCATTACCGATGTGACTTCAAAACTACTTACAG
 M A L L L E G G H Y R C D F K T T Y R

 610 620 630 640 650 660
 AGCTAAGAAGGTTGTCAGTTGCCAGACTATCATTTGTGGACCATCGCATTGAGATTGT
 A K K V V Q L P D Y H F V D H R I E I V

 670 680 690 700 710 720
 GAGCCACGACAAAGATTACAACAAGGTTAACGGCTTAAGCTGTATGAGCATGCCGAAGCTCATTCTGG
 S H D K D Y N K V K L Y E H A E A H S G

 730 740 750 760 770 780
 GCTGCCGAGGCAGGCCAAGTAAAGGCTTAATGAAAAGCCAAGACGACAACAAGGAGAAC
 L P R Q A K *

 790 800 810 820 830 840
 AAAGTATTTTTGTAAATTCAGGCATTTACTCGGAATTAGTATTGATACTTCG
 850 860 870 880 890 900
 ATTCAAGGATTGTTGGACTTGTAGAGACCAAGCTCTAGAGTTGTATTGTGAAAA
 910
 AAAGATAGTTCC 3'

(SEQ ID NOS: 25 & 26)

FIGURE 21

Green fluorescent protein homolog from *Montastraea annularis* mannFP (AY037766)

10 20 30 40 50 60
 5' TGGTTAACCGAGTCGGGGGGTCCCTGGCTAATAATTGATTCTATTGGGTGTGAC
 70 80 90 100 110 120
 ATTCAGGTTAACGAGCATCCTCAGTGCTGAGGCTCATTCACCCCTGGTGTGGAG
 130 140 150 160 170 180
 AGAGCAGATCGAGAACACCAAGAGCTTACGCTAAATCTTACCGCTCTACCACC
 190 200 210 220 230 240
 ATGAGTATGATTAAACCGAGAACATGAAGATCAAGATGGCTATGGACGGTGTAAACGGG
 M S M I K P E M K I K M R M D G A V N G

 250 260 270 280 290 300
 CACAAGTTCTGATTACAGGGAAAGGAAGCGCGAGCCTTCGAGGGAAAACAGACTATG
 H K F V I T G E G S G E P F E G K Q T M

 310 320 330 340 350 360
 AACCTGACAGTCATAGACGGCGGACCTCTGCCTTCGCTTCGACATCTTGACAACAGCA
 N L T V I D G G P L P F A F D I L T T A

 370 380 390 400 410 420
 TTTCGATTACGGCAMCAGGGTATTGCCAAATACCCAGAACAGACATCCCAGACTATTCAG
 F D Y G X R V F A K Y P E D I P D Y F K

 430 440 450 460 470 480
 CAGTCGTTCTGAGGGTTCTGGAACGAAGCATGACTTACGAAGACGGGGCATT
 Q S F P E G F S W E R S M T Y E D G G I

 490 500 510 520 530 540
 TGCATCGCACAATGACATAAAATGGAAGGCGACTGCTTTCCATATGAAATTGATTT
 C I A T N D I K M E G D C F S Y E I R F

 550 560 570 580 590 600
 GATGGGGTGAACCTTCCTGCCAATAGTCCAGTTATGCAAGAACCGTGAAATGGGAG
 D G V N F P A N S P V M Q K K T V K W E

 610 620 630 640 650 660
 CCATGCACTGRGGAAATGTATGTGCGTGATGGAGTGCTTAAAGGTGGCTTAAACATGGCT
 P C T X E M Y V R D G V L K G G L N M A

 670 680 690 700 710 720
 CTGTTGCTTGAAGGAGGTGGCATTCCGATGTGACTTGAAACTACTTACAAAGCTAAG
 L L L E G G H F R C D L K T T Y K A K

 730 740 750 760 770 780
 AAGGTTGCCAGATGCCAGACTATCACTTGTGAATCACCGACTTGAGATAACATGGCAT
 K V V Q M P D Y H F V N H R L E I T W H

 790 800 810 820 830 840
 GACGAGGATTACAACAATGTTAACGACTGCTGAGCATGCAAGCTCATTCTGGACTGCCA
 D E D Y N N V K L S E H A E A H S G L P

 850 860 870 880 890 900
 AGGCAGGCCAATAAAGGCTTGACGAAAGCCAAACGGCAAAGAGTACAAGAAAGTATA
 R Q A K *

 910 920 930 940 950 960
 TATAATGTATATTTCAACTGAAAGGCATTCCACTCGGAATTAGTATTGATACTTTC
 970 980 990 1000 1010 1020
 AATTCAAGGATTATTTGGGATTGCTAGCCACTAGCTTATTGTTAAATTAAAGTTAA
 1030 1040 1050 1060 1070 1080
 GACGGTTAGCATTTTCGGTATTACAACATAGGCACAGACGTCTAACCCAGTAGTG
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(SEQ ID NOS:27 & 28)

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							20		25				30		
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							35		40			45			
Gly	Pro	Leu	Pro	Phe	Ala	Phe	Asp	Ile	Leu	Ser	His	Ala	Phe	Arg	Tyr
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							65		70			75		80	
Lys	Gln	Ser	Leu	Pro	Glu	Gly	Phe	Thr	Trp	Glu	Arg	Val	Ser	Asn	Tyr
							85		90			95			
Glu	Asp	Gly	Gly	Val	Leu	Thr	Val	Lys	Gln	Glu	Thr	Ser	Leu	Glu	Gly
							100		105			110			

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 Thr Thr Tyr Lys Ser Lys Lys Gly Glu Pro Ala Lys Pro His Phe
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 Ile Thr Met Val Gly Gly Glu Phe Gln Tyr Asp Ile Arg Phe Asp Gly
 115 120 125
 Leu Asn Phe Pro Glu Asp Gly Pro Val Met Gln Lys Lys Thr Val Lys
 130 135 140
 Trp Glu Pro Ser Thr Glu Ile Met Tyr Met Gln Asn Gly Val Leu Lys
 145 150 155 160
 Gly Glu Val Asn Met Ala Leu Leu Gln Asp Lys Ser His Tyr Arg
 165 170 175
 Cys Asp Leu Lys Thr Thr Tyr Lys Ala Lys Asn Asn Val Pro His Pro
 180 185 190
 Pro Gly Tyr His Tyr Val Asp His Cys Ile Glu Ile Leu Glu Glu Arg
 195 200 205
 Lys Asp His Val Lys Leu Arg Glu His Ala Lys Ala Arg Ser Ser Leu
 210 215 220
 Ser Pro Thr Ser Ala Lys Glu Arg Lys Ala
 225 230

<210> 9
<211> 819
<212> DNA
<213> Scolymia cubensis

<400> 9
cctggtgatt tggacgagag cagatcgaga atagcaaggt tttaccagcg tgataattta 60
ctttacatct aacaacatgc aatctgctgg gaagaagaat gtcgttaagg acttcatgaa 120
gatcacactg cgtatggacg gtgctgtaaa cgggaagccc ttgcgggtta atggaacagg 180
agatggcaac ccttatggtg gaatacagag tttgaagctt accgtcgatg gcaacaaacc 240
tctgcctttt gcttttata tcttgcagc agcattccag tatggcaaca gggcattcac 300
cgaataccca aaagagatata cagactattt caagcagtcg ttttagtttgcgaggggtt 360
tacctggaa cgaagttca ctttgcaga cggggccatt tgcgtcgcca cgaacgatata 420
aaagatgggtt ggcgatgagt ttcaatataa cattcgattt gatgggtgtga atttccctga 480
agatggtccw gtyatgcaga agaaaaacggt gaagtgggag ccatccacag agataatgcg 540
tgtgcaaggt ggagtgcataa agggtgaggt taacatggct ctgttgctta aagacaaaag 600
ccattaccga tgcgtacttca aaactactta caaagctaag aatcctgtcc cgccgacggc 660
gcttccagac taccactatg tggatcactg tattgaaatc accgaggaaa atagggatta 720
cgttaagctg caggagttatg ctaaagctcg ttctggcctg cacctgccc aactgcaaaa 780
gtaaaggctt aggcgatagt caagacgaca acgagaaga 819

<210> 10
<211> 235
<212> PRT
<213> Scolymia cubensis

<400> 10
Met Gln Ser Ala Gly Lys Lys Asn Val Val Lys Asp Phe Met Lys Ile
 1 5 10 15
 Thr Leu Arg Met Asp Gly Ala Val Asn Gly Lys Pro Phe Ala Val Asn
 20 25 30
 Gly Thr Gly Asp Gly Asn Pro Tyr Gly Gly Ile Gln Ser Leu Lys Leu
 35 40 45
 Thr Val Asp Gly Asn Lys Pro Leu Pro Phe Ala Phe Asp Ile Leu Ser
 50 55 60
 Ala Ala Phe Gln Tyr Gly Asn Arg Ala Phe Thr Glu Tyr Pro Lys Glu
 65 70 75 80
 Ile Ser Asp Tyr Phe Lys Gln Ser Phe Glu Phe Gly Glu Gly Phe Thr
 85 90 95
 Trp Glu Arg Ser Phe Thr Phe Glu Asp Gly Ala Ile Cys Val Ala Thr
 100 105 110

Asn	Asp	Ile	Lys	Met	Val	Gly	Asp	Glu	Phe	Gln	Tyr	Asn	Ile	Arg	Phe
		115				120					125				
Asp	Gly	Val	Asn	Phe	Pro	Glu	Asp	Gly	Pro	Val	Met	Gln	Lys	Lys	Thr
		130				135					140				
Val	Lys	Trp	Glu	Pro	Ser	Thr	Glu	Ile	Met	Arg	Val	Gln	Gly	Gly	Val
					150					155					160
Leu	Lys	Gly	Glu	Val	Asn	Met	Ala	Leu	Leu	Leu	Lys	Asp	Lys	Ser	His
					165				170						175
Tyr	Arg	Cys	Asp	Phe	Lys	Thr	Thr	Tyr	Lys	Ala	Lys	Asn	Pro	Val	Pro
				180				185							190
Pro	Thr	Ala	Leu	Pro	Asp	Tyr	His	Tyr	Val	Asp	His	Cys	Ile	Glu	Ile
				195				200				205			
Thr	Glu	Glu	Asn	Arg	Asp	Tyr	Val	Lys	Leu	Gln	Glu	Tyr	Ala	Lys	Ala
					210			215			220				
Arg	Ser	Gly	Leu	His	Leu	Pro	Glu	Leu	Gln	Lys					
					225			230			235				

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<210> 11
<211> 807
<212> DNA
<213> Ricordea florida
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<210> 12
<211> 231
<212> PRT
<213> Ricordea florida

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<400> 12
Met Ser Ala Leu Lys Glu Glu Met Lys Ile Lys Leu Thr Leu Val Gly
  1           5           10           15
Val Val Asn Gly His Pro Phe Lys Ile Ile Gly Asp Gly Lys Gly Lys
  20          25          30
Pro Tyr Glu Gly Ser Gln Glu Leu Thr Leu Ala Val Val Glu Gly Gly
  35          40          45
Pro Leu Pro Phe Ser Tyr Asp Ile Leu Thr Thr Ile Val His Tyr Gly
  50          55          60
Asn Arg Ala Phe Val Asn Tyr Pro Lys Asp Ile Pro Asp Ile Phe Lys
  65          70          75          80
Gln Thr Cys Ser Gly Pro Gly Ala Gly Tyr Ser Trp Gln Arg Thr Met
  85          90          95
Ser Phe Glu Asp Gly Gly Val Cys Thr Ala Thr Ser His Ile Arg Val
 100         105         110
Asp Gly Asp Thr Phe Asn Tyr Asp Ile His Phe Met Gly Ala Asp Phe
 115         120         125

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Pro Leu Asn Gly Pro Val Met Gln Lys Arg Thr Val Lys Trp Glu Pro
 130 135 140
 Ser Thr Glu Ile Met Phe Gln Cys Asp Gly Leu Leu Arg Gly Asp Val
 145 150 155 160
 Ala Met Ser Leu Leu Leu Lys Gly Gly His Tyr Arg Cys Asp Phe
 165 170 175
 Lys Thr Ile Tyr Lys Pro Lys Lys Asn Val Lys Met Pro Gly Tyr His
 180 185 190
 Phe Val Asp His Cys Ile Glu Ile Thr Ser Gln Gln Asp Asp Tyr Asn
 195 200 205
 Val Val Glu Leu Tyr Glu Gly Ala Val Ala His Tyr Ser Pro Leu Gln
 210 215 220
 Lys Pro Cys Gln Ala Lys Ala
 225 230

<210> 13
<211> 796
<212> DNA
<213> Ricordea florida

<400> 13
 agtcacccctcg gtgttttttag gacaggaagg atcacgagca agagaagaac tgtgaaagtt 60
 aacactttac tctacttcta ccagcatgag tgcactcaa gaggaaatga aaatcaagct 120
 taaaatggtg ggcgttggta acgggcagtc attcagatc gatggggaa gaaaaggcaa 180
 accttacgag ggatcacaga aattaaccct tgaagtggtg gaaggagggc ctctgcttt 240
 ctcttatgat atcctgacaa cgatatttca gtatggcaac agggcattcg tgaactaccc 300
 aaaggacata ccagatattt tcaagcagac ctgctctggc cctgatggtg gattttctg 360
 gcaaaggacc atgactttag aagacggagg gtttgact gcttcaaacc acatcagcgt 420
 ggacggcgcac actttttat atgtgataag atttaatgga gagaattttc ctccaaatgg 480
 tccagtaatg cagaaaagaa cagtgaaatg ggagccatcc actgagataa tgtttgaacg 540
 tgatggattg ctgagggggt acattgccat gtctctgtt ctgaaaggag gcggccattt 600
 ccgatgtgac ttataaacta tttatacacc caagaggaag gtcaacatgc caggttacca 660
 ttttggcact cactgcattt agatacagaa gcacgacaag gattacaaca tggctgtgct 720
 ctctgaggat gctgttagccc acaactctcc tctggagaaaa aaaagccaag caaaggcgta 780
 aagccaaaca acctaa 796

<210> 14
<211> 231
<212> PRT
<213> Ricordea florida

<400> 14
 Met Ser Ala Leu Lys Glu Glu Met Lys Ile Lys Leu Lys Met Val Gly
 1 5 10 15
 Val Val Asn Gly Gln Ser Phe Gln Ile Asp Gly Glu Gly Lys Gly Lys
 20 25 30
 Pro Tyr Glu Gly Ser Gln Lys Leu Thr Leu Glu Val Val Glu Gly Gly
 35 40 45
 Pro Leu Leu Phe Ser Tyr Asp Ile Leu Thr Thr Ile Phe Gln Tyr Gly
 50 55 60
 Asn Arg Ala Phe Val Asn Tyr Pro Lys Asp Ile Pro Asp Ile Phe Lys
 65 70 75 80
 Gln Thr Cys Ser Gly Pro Asp Gly Gly Phe Ser Trp Gln Arg Thr Met
 85 90 95
 Thr Tyr Glu Asp Gly Gly Val Cys Thr Ala Ser Asn His Ile Ser Val
 100 105 110
 Asp Gly Asp Thr Phe Tyr Tyr Val Ile Arg Phe Asn Gly Glu Asn Phe
 115 120 125
 Pro Pro Asn Gly Pro Val Met Gln Lys Arg Thr Val Lys Trp Glu Pro
 130 135 140

Ser Thr Glu Ile Met Phe Glu Arg Asp Gly Leu Leu Arg Gly Asp Ile
 145 150 155 160
 Ala Met Ser Leu Leu Leu Lys Gly Gly His Tyr Arg Cys Asp Phe
 165 170 175
 Lys Thr Ile Tyr Thr Pro Lys Arg Lys Val Asn Met Pro Gly Tyr His
 180 185 190
 Phe Val Asp His Cys Ile Glu Ile Gln Lys His Asp Lys Asp Tyr Asn
 195 200 205
 Met Ala Val Leu Ser Glu Asp Ala Val Ala His Asn Ser Pro Leu Glu
 210 215 220
 Lys Lys Ser Gln Ala Lys Ala
 225 230

<210> 15
 <211> 795
 <212> DNA
 <213> Montastraea cavernosa

<400> 15
 acgcaggat tcaccctgg gatttggaaag agagcagacc gagaacaaca agagctgtat 60
 aaggctgata tcttacttta cgtctaccat catgagtgtg attaaatcag tcatgaagat 120
 caagctcggt atggaaggca gtgttaaacgg gcacaacttc gtaattgtt gagaaggaga 180
 aggcaagcct tatgagggaa cacagagtat ggaccttaca gtcaaagaag ggcacccct 240
 gccttcgccc tacgatatac tgacaacagt attccattac ggcaataggg tattcgcaaa 300
 atacccaaaa catatccca gactattcaa gcagatgtt cctgaggagt attcctggga 360
 acgaagcatg aatttcgaag gcggggcat ttgcaccgccc aggaacgaga taacaatgg 420
 aggccgactgt ttttcaata aagttcgatt tgatgggttg aacttcccccc ccaatggtcc 480
 agtcatgcag aagaagacgc taaaaatggg gccatccact gaaaaaaatgt atgtgcgtga 540
 tggagtgctg acgggtgata tcaacatggc tttgttgctt gaaggaggtg gccattaccg 600
 atgtgacttc agaactactt acagagctaa gaagaagggt gtcaagttac cagattatca 660
 ctttgaggat cactccattt agattttgcg ccatgacaaaa gaatacactg aggttaagct 720
 gtatgagcat gcccgaagctc attctggct gcccgggtg gcaaaagtaaa ggcttaacga 780
 aaagccaaga ccaca 795

<210> 16
 <211> 235
 <212> PRT
 <213> Montastraea cavernosa

<400> 16
 Arg Leu Ile Ser Tyr Phe Thr Ser Thr Ile Met Ser Val Ile Lys Ser
 1 5 10 15
 Val Met Lys Ile Lys Leu Arg Met Glu Gly Ser Val Asn Gly His Asn
 20 25 30
 Phe Val Ile Val Gly Glu Gly Glu Lys Pro Tyr Glu Gly Thr Gln
 35 40 45
 Ser Met Asp Leu Thr Val Lys Glu Gly Ala Pro Leu Pro Phe Ala Tyr
 50 55 60
 Asp Ile Met Thr Thr Val Phe His Tyr Gly Asn Arg Val Phe Ala Lys
 65 70 75 80
 Tyr Pro Lys His Ile Pro Asp Tyr Phe Lys Gln Met Phe Pro Glu Glu
 85 90 95
 Tyr Ser Trp Glu Arg Ser Met Asn Phe Glu Gly Gly Ile Cys Thr
 100 105 110
 Ala Arg Asn Glu Ile Thr Met Glu Gly Asp Cys Phe Phe Asn Lys Val
 115 120 125
 Arg Phe Asp Gly Val Asn Phe Pro Pro Asn Gly Pro Val Met Gln Lys
 130 135 140
 Lys Thr Leu Lys Trp Glu Pro Ser Thr Glu Lys Met Tyr Val Arg Asp
 145 150 155 160

Gly Val Leu Thr Gly Asp Ile Asn Met Ala Leu Leu Leu Glu Gly Gly
 165 170 175
 Gly His Tyr Arg Cys Asp Phe Arg Thr Thr Tyr Arg Ala Lys Lys Lys
 180 185 190
 Gly Val Lys Leu Pro Asp Tyr His Phe Glu Asp His Ser Ile Glu Ile
 195 200 205
 Leu Arg His Asp Lys Glu Tyr Thr Glu Val Lys Leu Tyr Glu His Ala
 210 215 220
 Glu Ala His Ser Gly Leu Pro Arg Val Ala Lys
 225 230 235

<210> 17
 <211> 1066
 <212> DNA
 <213> Montastraea cavernosa

<400> 17
 attcgccctg gtgatttgg aagagacaga tcgagaacaa caagagctgt aagggttata 60
 tcttacttac gtctaccatc atgacaagtg ttgcacagga aaagggtgtg attaaaccag 120
 acatgaagat gaagctgcgt atggaagggtg ctgtaaacgg gcacaagttc gtgggtgaag 180
 gagatggaaa agggaaaggct ttcgacggaa cacagacttggacattaca gtcatagaag 240
 ggcgcaccatt gccttcgct tacgatatct tgacaacagt attcgattac ggcaacagg 300
 tattcgccaa atacccagaa gacatagcag attatttcaa gcagacgtt cctgagggg 360
 acttctggaa acgaagcatg acatacgaag accagggtt ttgcatcgcc acaaacgaca 420
 taacaatgat ggaaggcggtc gacgactgtt ttgcctataa aattcgattt gatgggtgt 480
 actttcttcgcaatggtcca gttatgcaga ggaagacgct gaaatgggag ccatccactg 540
 agataatgtatgcgtgat ggagtgtga agggtgatgt taacatggct ctgttgctt 600
 aaggaggtgg ccattaccga tgtgacttca aaactactta caaagctaag aaggttgc 660
 ggttgcaga ctatcactt gtggaccatc gcattggatgtt gatggccac gacaaagatt 720
 acaacaaggtaaagctgcac gaggatggccaa gaaatgggag ccatccactg 780
 agtaaaggct taatgaaaatg tcaagacgac aacgaggaga aacaaatgtt tttttgtt 840
 aatttgaagg catttactcg gaatttagtat ttgataacttt cgattcaagg atttgttcc 900
 ggattttgtt aagacttagctt gatggccac gacaaagattt ttccagttt 960
 tgcgggattt cagcatgggg atagactttt taaactcaatgtt gatggccac 1020
 aaaactgttagt tgagaataaa ctgttatcg aagccgaaaa aaaaaaa 1066

<210> 18
 <211> 234
 <212> PRT
 <213> Montastraea cavernosa

<400> 18
 Met Thr Ser Val Ala Gln Glu Lys Gly Val Ile Lys Pro Asp Met Lys
 1 5 10 15
 Met Lys Leu Arg Met Glu Gly Ala Val Asn Gly His Lys Phe Val Val
 20 25 30
 Glu Gly Asp Gly Lys Gly Lys Pro Phe Asp Gly Thr Gln Thr Met Asp
 35 40 45
 Leu Thr Val Ile Glu Gly Ala Pro Leu Pro Phe Ala Tyr Asp Ile Leu
 50 55 60
 Thr Thr Val Phe Asp Tyr Gly Asn Arg Val Phe Ala Lys Tyr Pro Glu
 65 70 75 80
 Asp Ile Ala Asp Tyr Phe Lys Gln Thr Phe Pro Glu Gly Tyr Phe Trp
 85 90 95
 Glu Arg Ser Met Thr Tyr Glu Asp Gln Gly Ile Cys Ile Ala Thr Asn
 100 105 110
 Asp Ile Thr Met Met Glu Gly Val Asp Asp Cys Phe Ala Tyr Lys Ile
 115 120 125
 Arg Phe Asp Gly Val Asn Phe Pro Ala Asn Gly Pro Val Met Gln Arg
 130 135 140

Lys Thr Leu Lys Trp Glu Pro Ser Thr Glu Ile Met Tyr Ala Arg Asp
 145 150 155 160
 Gly Val Leu Lys Gly Asp Val Asn Met Ala Leu Leu Leu Glu Gly Gly
 165 170 175
 Gly His Tyr Arg Cys Asp Phe Lys Thr Thr Tyr Lys Ala Lys Lys Val
 180 185 190
 Val Arg Leu Pro Asp Tyr His Phe Val Asp His Arg Ile Glu Ile Val
 195 200 205
 Ser His Asp Lys Asp Tyr Asn Lys Val Lys Leu His Glu His Ala Glu
 210 215 220
 Ala Arg His Gly Leu Ser Arg Lys Ala Lys
 225 230

<210> 19
 <211> 898
 <212> DNA
 <213> *Condylactis gigantea*

<400> 19
 acagctttc atccacgctc attcaagacg ccgtcaactt tattccagtc agaaaaatgt 60
 atccttggat caaggaaacc atgcgcagta aggtttacat ggaaggagat gttacaacc 120
 acgccttcaa gtgcactgca gtaggagaag gaaaaccata caaaggctca caagacctga 180
 cgattaccgt cactgaagga ggtcctctgc catttgcttt cgacattctt tcacacgcct 240
 ttcagtagg caacaagggtt ttcaccgatt accccgacga tattcctgat ttctttaagc 300
 agtctcttc ggtatggttt acttgagaa gagaaggac statgacgat ggaggagtc 360
 tcacagttac ccaagacact agtctgaagg gagattgcat tatttgcac attaaagtcc 420
 atggcactaa cttcccccgaa aatggtccgg tgatgcaaaa caagaccgat ggatgggagc 480
 catccagcac tgaaacgggtt attccacaag atggaggaat tggtgctgcg cgatcacccg 540
 cactaaggct gctgtataaa ggtcatctta tctgccacat gggaaacaact tacaaggccaa 600
 acaaagaggt gaagctgcca gaactccact ttcatcattt gcgaatggaa aagctgagtg 660
 ttagtgacga tgggaagacc attaagcgc acgagtagt ggtggctagc tactccaaag 720
 tgccttcgaa gataggacgt caatgatcat ttcccttatt aaatatcaat gatgtggctt 780
 tcaattttcc aaaattttgt taagacatag gtctttgga tttttgtaa ccccaacctt 840
 aattcccaat aatttttgtt ggaaagtcaa ataaaaccag ccttccctgg gcctttaa 898

<210> 20
 <211> 229
 <212> PRT
 <213> *Condylactis gigantea*

<400> 20
 Met Tyr Pro Trp Ile Lys Glu Thr Met Arg Ser Lys Val Tyr Met Glu
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 Gly Asp Val Asn Asn His Ala Phe Lys Cys Thr Ala Val Gly Glu Gly
 20 25 30
 Lys Pro Tyr Lys Gly Ser Gln Asp Leu Thr Ile Thr Val Thr Glu Gly
 35 40 45
 Gly Pro Leu Pro Phe Ala Phe Asp Ile Leu Ser His Ala Phe Gln Tyr
 50 55 60
 Gly Asn Lys Val Phe Thr Asp Tyr Pro Asp Asp Ile Pro Asp Phe Phe
 65 70 75 80
 Lys Gln Ser Leu Ser Asp Gly Phe Thr Trp Arg Arg Val Ser Thr Tyr
 85 90 95
 Asp Asp Gly Gly Val Leu Thr Val Thr Gln Asp Thr Ser Leu Lys Gly
 100 105 110
 Asp Cys Ile Ile Cys Asn Ile Lys Val His Gly Thr Asn Phe Pro Glu
 115 120 125
 Asn Gly Pro Val Met Gln Asn Lys Thr Asp Gly Trp Glu Pro Ser Ser
 130 135 140
 Thr Glu Thr Val Ile Pro Gln Asp Gly Ile Val Ala Ala Arg Ser

145	150	155	160
Pro Ala Leu Arg Leu Arg Asp Lys Gly His	Leu Ile Cys His	Met Glu	
165	170	175	
Thr Thr Tyr Lys Pro Asn Lys Glu Val Lys	Leu Pro Glu Leu His	Phe	
180	185	190	
His His Leu Arg Met Glu Lys Leu Ser Val	Ser Asp Asp Gly Lys	Thr	
195	200	205	
Ile Lys Gln His Glu Tyr Val Val Ala Ser	Tyr Ser Lys Val Pro	Ser	
210	215	220	
Lys Ile Gly Arg Gln			
225			

<210> 21
<211> 1030
<212> DNA
<213> Agaricia fragilis

<400> 21
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tttctactct aaagaacgtc atcatcatcg ttattatata ctcctgcagc acttgtgctg 120
tttggtcgaa ttcaaactct gaatcttctt tcactaatgg gattgcagag gaaatgaaga 180
ctagggtaca ttggaggtt actgttaacg ggcactcctt tacaattaaa ggcgaaggaa 240
gaggctaccc ttacaaagga gaacagttt tgagccttga ggtcgtcaat ggtgctcctc 300
tgccgttctc ttttgatattc ttgacaccag catttatgtt tggcaacaga gtgttcacca 360
agtaccaccc aaacataccca gactattca agcagacgtt tccgttggg tatactggg 420
aaagaaacat tcccttggaa gatcaggccg cgtgcacggg aaccagccac ataagatgg 480
aagaggaaga gaggcggtt gtaataacg tcagatttca ctgtgtgaac tttcccccta 540
atggtccagt catgcagagg aggatactga aatgggagcc atccactgaa aacatttatac 600
cgcgtgatgg gtttctggag ggccatgttgc atatgacttc tcgggttggaa ggagggtggct 660
attaccgagc ttagttcaaa agtacttaca aagggaaagac cccagtcgc gacatgccag 720
actttcactt catagaccac cgcatggaga ttacggagca tgacgaagac tacaccaatg 780
ttgagctgca tgacgtatcc tgggctcggtt actctatytgc gccgactatg taagcggaaa 840
aggcaaggca acaagacgca aaaccggccct gtttgcctt tttcataaga gatttgacaa 900
ccgtgggttct ttgccatttta atttgaatttta gtttaaatttta aatctttggg attgatgttag 960
acgctttgggt tgctaagttaa gaaaacattt tgatttgc ctgaagcaaa 1020
aaaaaaaaaa 1030

<210> 22
<211> 259
<212> PRT
<213> Agaricia fragilis

<400> 22
Met Ala Ile Ser Thr Leu Lys Asn Val Ile Ile Ile Val Ile Ile Tyr
1 5 10 15
Ser Cys Ser Thr Cys Ala Val Trp Ser Asn Ser Asn Ser Glu Ser Ser
20 25 30
Phe Thr Asn Gly Ile Ala Glu Glu Met Lys Thr Arg Val His Leu Glu
35 40 45
Gly Thr Val Asn Gly His Ser Phe Thr Ile Lys Gly Glu Gly Arg Gly
50 55 60
Tyr Pro Tyr Lys Gly Glu Gln Phe Met Ser Leu Glu Val Val Asn Gly
65 70 75 80
Ala Pro Leu Pro Phe Ser Phe Asp Ile Leu Thr Pro Ala Phe Met Tyr
85 90 95
Gly Asn Arg Val Phe Thr Lys Tyr Pro Pro Asn Ile Pro Asp Tyr Phe
100 105 110
Lys Gln Thr Phe Pro Glu Gly Tyr His Trp Glu Arg Asn Ile Pro Phe
115 120 125
Glu Asp Gln Ala Ala Cys Thr Val Thr Ser His Ile Arg Leu Glu Glu

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<210> 23
<211> 1024
<212> DNA
<213> Ricordea florida
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<400> 23
agccacttcg gtgtcttgc gagaggaagg atcacgaaca agagaagac tggaaaagg 60
aaaattttac ttacttctt ccagcatgaa tgcacttcaa gaggaaatga aatcaagct 120
tacaatggtg ggcgttgtta acgggcagtc atttaagatc gatgggaaag gaaaaggaa 180
accttacgag ggatcacagg aattgaccct taaaagtggtg gaaggcgggc ctctgcttt 240
ctcttatgat atcctgacaa cgatatttca gtatggcaac agggcattcg tgaactaccc 300
aaaggacata ccagatattt tcaagcaaac gtgttctggt cttgatggcg gatattcg 360
gcaaaggacc atgactttatg aggacggagg ggtttgtact gctacaagca acgtcagcgt 420
ggtcggcgac actttcaatt atgaaattca ctttatgggg gcgaattttc ctccaaatgg 480
tccrqtatg cagaaaagaa cagtgaagtgg gggcccttc actgagataa tggttgaacg 540
tgatggattt ctgggggttg atgttccat gtctctgttg ctgaaaaggag gcgaccattt 600
ccgatgtgac ttaaaacta ttataaacc caacaagaag gtcaagctgc caggttacca 660
ttttgtggac cactgcattt agataaaagag tcaagagaat gattacaaca tggttgcgt 720
cttgaggat gctgttagcac actactctcc tctggggaaa aagagccagg caaaggcgt 780
aatccaaaca acctaagaag acgacaaggc attcaatcta atcgatgtt tgaattttt 840
gttaggaatg tgggggtca gacttaggtct agaacgttc atttggctg gatttggttt 900
actcaggat agacaagaaa aaaactttaa atgacttggg ttggatttag ctttcggcac 960
tgtcaattcc ggattcctta gaaatatttgc agaccaagcc ttttttgag ctgagaacgt 1020
aattc 1024

<210> 24
<211> 231
<212> PRT
<213> Ricordea florida

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<400> 24
Met Asn Ala Leu Gln Glu Glu Met Lys Ile Lys Leu Thr Met Val Gly
  1           5           10          15
Val Val Asn Gly Gln Ser Phe Lys Ile Asp Gly Lys Gly Lys Gly Lys
  20          25          30
Pro Tyr Glu Gly Ser Gln Glu Leu Thr Leu Lys Val Val Glu Gly Gly
  35          40          45
Pro Leu Leu Phe Ser Tyr Asp Ile Leu Thr Thr Ile Phe Gln Tyr Gly
  50          55          60
Asn Arg Ala Phe Val Asn Tyr Pro Lys Asp Ile Pro Asp Ile Phe Lys
 65          70          75          80
Gln Thr Cys Ser Gly Leu Asp Gly Gly Tyr Ser Trp Gln Arg Thr Met

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	85		90		95										
Thr	Tyr	Glu	Asp	Gly	Gly	Val	Cys	Thr	Ala	Thr	Ser	Asn	Val	Ser	Val
		100						105						110	
Val	Gly	Asp	Thr	Phe	Asn	Tyr	Glu	Ile	His	Phe	Met	Gly	Ala	Asn	Phe
		115						120				125			
Pro	Pro	Asn	Gly	Pro	Val	Met	Gln	Lys	Arg	Thr	Val	Lys	Trp	Glu	Pro
		130					135				140				
Ser	Thr	Glu	Ile	Met	Phe	Glu	Arg	Asp	Gly	Leu	Leu	Arg	Gly	Asp	Val
		145				150				155				160	
Pro	Met	Ser	Leu	Leu	Leu	Lys	Gly	Gly	Asp	His	Tyr	Arg	Cys	Asp	Phe
		165					170					175			
Lys	Thr	Ile	Tyr	Lys	Pro	Asn	Lys	Lys	Val	Lys	Leu	Pro	Gly	Tyr	His
		180					185					190			
Phe	Val	Asp	His	Cys	Ile	Glu	Ile	Lys	Ser	Gln	Glu	Asn	Asp	Tyr	Asn
		195					200					205			
Met	Val	Ala	Leu	Phe	Glu	Asp	Ala	Val	Ala	His	Tyr	Ser	Pro	Leu	Glu
		210				215					220				
Lys	Lys	Ser	Gln	Ala	Lys	Ala									
							225					230			

<210> 25

<211> 913

<212> DNA

<213> Montastraea cavernosa

<400> 25

agagctgttag ggtgatatct tacttacgtc taccatcatg accagtgttg cacagggaaaa 60
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 caagttcgtg attgaaggag atggaaaagg gaagcctttc gacggaacac agactatgg 180
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<211> 234

<212> PRT

<213> Montastraea cavernosa

<400> 26

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Gly	Val	Leu	Lys	Gly	Asp	Val	Asn	Met	Ala	Leu	Leu	Leu	Glu	Gly	Gly
			165			170							175		
Gly	His	Tyr	Arg	Cys	Asp	Phe	Lys	Thr	Thr	Tyr	Arg	Ala	Lys	Lys	Val
			180			185							190		
Val	Gln	Leu	Pro	Asp	Tyr	His	Phe	Val	Asp	His	Arg	Ile	Glu	Ile	Val
			195			200							205		
Ser	His	Asp	Lys	Asp	Tyr	Asn	Lys	Val	Lys	Leu	Tyr	Glu	His	Ala	Glu
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<210> 27

<211> 1133

<212> DNA

<213> Montastraea annularis

<400> 27

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<210> 28

<211> 224

<212> PRT

<213> Montastraea annularis

<220>

<221> VARIANT

<222> 65, 144

<223> Xaa = Any Amino Acid

<400> 28

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Xaa	Arg	Val	Phe	Ala	Lys	Tyr	Pro	Glu	Asp	Ile	Pro	Asp	Tyr	Phe	Lys
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Gln	Ser	Phe	Pro	Glu	Gly	Phe	Ser	Trp	Glu	Arg	Ser	Met	Thr	Tyr	Glu
					85				90				95		
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			100					105					110		
Cys	Phe	Ser	Tyr	Glu	Ile	Arg	Phe	Asp	Gly	Val	Asn	Phe	Pro	Ala	Asn
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						165			170				175		
Tyr	Lys	Ala	Lys	Lys	Val	Val	Gln	Met	Pro	Asp	Tyr	His	Phe	Val	Asn
							180		185			190			
His	Arg	Leu	Glu	Ile	Thr	Trp	His	Asp	Glu	Asp	Tyr	Asn	Asn	Val	Lys
					195			200				205			
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					210			215				220			